

OM of: US-09-303-518D-125 to: A.Geneseq.032802.* out_format: pfs
Date: Jun 30, 2002 6:38 AM

About: Results were produced by the GenCore software, version 4.5,
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Command line parameters:

-MODE=frame+np2 model -DEV=xlh
-o=/gen2.1/USPbio/US09303518/runat_28062002_142712_4291/app-query.fasta.1.23501
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-GAPEXT=4.000 -MINMATCH=0.100 -LOOPEL=0.000 -LOOPEXT=0.000
-GAPOP=6.000 -GAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500
-GAPOP=6.000 -GAPEXT=7.000 -XGAPOP=10.000 -XGAPEXT=0.500
-DELOP=6.000 -DELEXT=7.000 -START=1 -MATRIX=blcsu6m2
-TRANS=human40.cdt -LIST=100 -DOCALIGN=200 -THR_SCORE=pct
-THR_MAX=100 -THR_MIN=0 -ALIGN=45 -MODE=LOCAL -OUTFMT=pfs
-NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000
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-DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -NO_XLPXY -WAIT -THREADS=1

Search information block:

Query: US-09-303-518D-125
Query length: 1344
Database: A.Geneseq.032802.*
Database sequences: 74574
Database length: 11073796
Search time (sec): 627.340000

score_list:

Seqence Strd Orig ZScore EScore Len i Documentation

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ID: AAV38561 standard; Protein: 447 AA.

AAV38561;
08-Oct-1999 (first entry)
Neisseria meningitidis antigen encoded by ORF22.
Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;

KM treatment; Neisseria infection; meningitis; septicaemia; gonorrhea.
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 OS Neisseria meningitidis.
 PN W09924578-A2.
 XX
 PD 20-MAY-1999.
 XX
 PF 09-OCT-1998; 98MO-IB01665.
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 PR 01-SEP-1998; 98GB-0019016.
 PR 06-NOV-1997; 97GB-0023516.
 PR 14-NOV-1997; 97GB-0024190.
 PR 18-NOV-1997; 97GB-0024386.
 PR 27-NOV-1997; 97GB-0025158.
 PR 10-DEC-1997; 97GB-0026147.
 PR 14-JAN-1998; 98GB-0000759.
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 PA (CHIR-) CHIRON SPA.
 XX
 PI Grandi G, Masignani V, Pizzo M, Rappuoli R, Scarlato V;
 XX WPI: 1999-327407/27.
 DR N-PSDB; AAZ12026.
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 PT Proteins from Neisseria meningitidis and N. gonorrhoeae useful for
 diagnosis, treatment and prevention of infection
 XX
 PS Claim 4; Page 123; 524pp; English.
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 CC Amino acid sequences AAY38499-Y38944 represent Neisseria meningitidis
 CC and N. gonorrhoeae antigenic proteins. They are encoded by open
 CC reading frames (ORFs) AAZ1972-Z12358. The antigenic proteins,
 CC their fragments, their nucleic acids and antibodies are used for
 CC diagnosis, prevention (as vaccines) or treatment of Neisseria
 CC infections, such as meningitis, septicaemia and gonorrhea. Both
 CC organisms are closely related. Fragments of the nucleic acids
 CC are useful as hybridisation probes and antisense reagents.
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 SQ Sequence 447 AA;

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 Quality: 2289.00 Gaps: 0
 Ratio: 5.121 Percent Identity: 100.000
 Percent Similarity: 100.000

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 US-09-303-518D-125 x AAY38561 ..
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 17 uGlnAlaValIyrAspGlyProAlaIleThrGlnValAlaLeuLeuGlyg 34
 101 AAGAAATATGCGGATGCGCCCTCGATGAAGTCAAGGAAGCGGATGCC 150
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 151 GTCAAAAAAGCCAAAGTCTGTTGAAGACAAAAGAATCCGGCGGTGT 200
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 251 AGCGGTAAGTCAAGTCAAGTCAAGTCAAGTCAAGTCAAGTCAAGTCA 300

84 ysArgValLeuGlnSerValIleAlaValGlyGlyAsnAspGluIle 100
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 101 GluPheGluArgTyAlaProGlnAlaLeuAlaAsnLeuSerGlyGluI 117
 351 AGTGCCTGCAACCTGATCCAAATCCGGTGTGAGCTGCGTGCACCC 400
 117 uValAlaArgAsnLeuIleGlnSerGlyLeuThrAlaLeuArgThrAl 134
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 134 rgProPheSerLysIleProAlaValAlaSpAlaGluProPheAlaIlePhe 150
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 151 ValAsnAlaMetAspThrAsnProLeuAlaAlaAspProThrValIleI 167
 501 CAAGAAGCCGCGGAGATTCAAACGCGCGCTGTGATTAAGCCGTT 550
 167 elySGlAlaIleAlaGluAspPheLysArgGlyLeuLeuValLeuSerArgL 184
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 184 eutHrGluArgLysIleHisValLysAlaAlaGlyAlaAspValPro 200
 601 TCTGAATAATGCTGCCACATCGAAGCACATGAATTCGGCGCGCCATCC 650
 201 SerGluAsnAlaIleAsnIleGluThrHisGluPheGlyProHisPr 217
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 251 LeuPheAlaIleThrGlyArgLeuAsnThrGluArgValIleAlaLeuGlyG 267
 801 TTCTCAAGTCAACCAACCGCGCTTGGCTACCGCTTTGGTGGCAAG 850
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 367 aValAsnGlyGlyAspArgAlaMetValProIleGlyThrTyGluArgV 384
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401 GlysPThrAspSerAlaGlnAlaLeuGlyCysLeuGlnLeuAspGlnG 417
1251 AGACCTCGCTTGTGACAGCTCTGCTGCGCCGGCAATACGAATACGCGC 1300
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seq_documentation_block:
ID AAV38564 standard; Protein; 447 AA.
AC AAV38564;
XX
XX 08-OCT-1999 (first entry)
XX
XX Neisseria gonorrhoeae antigen encoded by ORF22.
XX
XX Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
KM treatment; Neisseria infection; meningitis; septicemia; gonorrhea.
XX
XX Neisseria gonorrhoeae.
XX
XX WO924578-A2.
XX
XX 20-MAY-1999.
XX
XX 09-OCT-1998; 98MO-IB01665.
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XX 01-SEP-1998; 98GB-0019016.
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XX 14-NOV-1997; 97GB-0024190.
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XX 18-NOV-1997; 97GB-0024386.
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XX 27-NOV-1997; 97GB-0025158.
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XX PI Grandi G, Masignani V, Pizza M, Rappuoli R, Scarlato V;
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XX WPI; 1999-327407/27.
XX
XX DR N-PSDB; AA212028.
XX
XX Proteins from Neisseria meningitidis and N. gonorrhoeae useful for
PT diagnosis, treatment and prevention of infection
XX
XX
XX Claim 4; Page 125; 524pp; English.
XX
XX
XX Amino acid sequences AAV38499-Y38944 represent Neisseria meningitidis
CC and N. gonorrhoeae antigenic proteins. They are encoded by open
CC reading frames (ORFs) AA211972-212358. The antigenic proteins,
CC their fragments, their nucleic acids and antibodies are used for
CC diagnosis, prevention (as vaccines) or treatment of Neisseria
CC infections, such as meningitis, septicemia and gonorrhea. Both
CC organisms are closely related. Fragments of the nucleic acids
CC are useful as hybridisation probes and antisense reagents.
XX
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XX Sequence 447 AA;

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Ratio: 4.980 Gaps: 0
Percent Similarity: 99.776 Percent Identity: 96.197

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17 uGlnValIleTyrAspGlyProAlaIleThrGlnValAlaLeuLeuGlyG 34
101 AAGCATATGCGGCTATCGCCCGCTCGATGAAGTGAAGGACGCGATGCC 150
34 IuGlnTyrValGlyMetArgProSerMetLysIleLysGlnGlyAla 50
151 GTCAAAAAAGGCCAAGTCTGTTGAAGACAAAAAGAAATCCGGCGTGT 200
51 ValLysLysGlyGlnValLeuPheGlnAspLysLysAsnProGlyValVa 67
201 GTTACTGCGCGCGCTTCAGCAAAATCGCCGATTCACCTGGCGAA 250
67 IPhetnAlaProAlaSerGlyLysIleAlaIleHisArgGlyGlu 84
251 AGCGCTACTCTCAGTCAGTCGATTCGCGTTCGAAGCGCAACGACAAATC 300
84 ysArgValLeuGlnSerValIleAlaValGlnLysAsnSpGluIle 100
301 GAGTTTGAACGCTACGCACTCGAAGCGCTGGCAAACTTAAGCGCGAAGA 350
101 GluPheGlnArgTyrValProGlnAlaLeuAlaLysLeuSerSerGln 117
351 AGTCGCGCGCAACCTGATCCGCTTGGATCGATCGCGCGACCC 400
117 sValArgArgAsnLeuIleGlnSerGlyLeuTyrPheAlaLeuArgTha 134
401 GTCCGTTACGCAAAATTCCTGCGCTGATGCCGAGCGCTTCGCCATCTTC 450
134 rGProheSerLysIleProAlaValAlaSpAlaGlnProheAlaIlePhe 150
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501 CAAGAAGCGCGGAGGATTCGAAAGCGCGCTGATGAGCGCTT 550
167 eLysGlnAlaAlaGlnAspPheLysArgLysLeuValLeuSerArgL 184
551 TGACCGACGCAAAATCATGTTTGAAGCGAGCTGCGCAGACGTGCCG 600
184 eThrGlnArgLysIleHisValCysLysAlaIleAlaLysAlaSpValPro 200
601 TCTGAAAATGCTCCCAACATCGAACACATGAATTCGGCGCCGCGATCC 650
201 SerGlnAsnAlaAlaAsnIleGlnThrHisGlnPheGlyGlyProHisPr 217
651 TGCCGGTTGAGTGGACGACATTCATTCATTCGACGCGCGGCGGCA 700
217 oAlaGlyLeuSerGlyThrHisIleHisPheIleGlnProValGlyAla 234
701 ATAAACCGTGTGACATCAATTCATCAAGATGTAATTCATTCGCGCT 750
234 snLysThrValTrpThrIleAsnTyrGlnAspAlaIleAlaIleGlyArg 250
751 TTGTTTGAACAGCGCGCTCTGAACACCGAGCGGTGATTCCTAGAGTG 800
251 LeuPheValThrGlyArgLeuAsnThrGlnArgValAlaLeuGlyG 267
801 TTCTCAAGTCAACAACCGCGCTTCGGGTACGTTTGGGTGGCGCAAG 850
267 yLeuGlnValAsnLysProArgLeuLeuArgThrValLeuGlyAlaLysV 284

```

851 TATGCAAAATACGCGGGCGAATTGGTTGACACAGACACCGCGTGATT 900
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
284 aIserscinleuthralaglyluleuValaspAlaspasnargValile 300
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
301 SeeglySerValleuasnlglyAlaleaglnglyAlahlsAspPtyrle 317
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
951 GGGAGCGCTACCAATCAAGATTCCGTTATCGAAGAGCGCGCACAAAG 1000
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
317 uGlymrGlyrHlsAsnglnlleserValilegluclymrGlyrGlysg 334
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
1001 ACCTGTTGGCGTGGTGGCGCGACCGGACAAATCTCCATCCAGCGGT 1050
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
334 luleuphegllyrPylAlaProglInProasplyrtySerlethrarg 350
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
1051 ACAACCTCGGCGCATTTCTCTGAAAAACAACCTTCAAGTTCAACACAGC 1100
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351 ThrThrleuGlyHlsPheleuLysasnLysleuPheLysPheThrThrAl 367
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1101 CGTCAACGGCGCGACCGCGCATGTGCGCATTTGTTACTTACGAGCGG 1150
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367 aValasnlglylaspargAlaMetValProlellyThrlyrGlymrGly 384
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
1151 TGATGCCCTTGATATCTGCGCCACCTGCTTTTGGCGGATTAACTGTC 1200
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384 alMetProleuaspIleleuProThrleuLeuAlargAspleuIleVal 400
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1201 GCGGATACCGACAGCGCGCGCATTTGGTGTCTTGAATTGGACGAAGA 1250
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401 GlyaSPThrAspSerAlaGlnAlaLeuGlyCysleuGluLeuAspGluGl 417
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
1251 AGACCTCGCTTGTGACGCTTGTGCTGCGCGCAATACGAATACGCGCC 1300
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417 uAspleuAlaLeuGlySerPheValCysProgllyrtyrGlymrGlyr 434
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
1301 CGCTGTGGCGCAAGTCTGCAAAACCATTTGAAGAGGAAGGC 1341
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
434 rleuLeuAlarglyValleuGluThrIleGlymrGlymrGlymr 447
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seq_documentation_block:
ID AAV38562 standard; Protein; 447 AA.
XX
AC AAV38562;
XX
DT 08-OCT-1999 (first entry)
XX
DE Neisseria meningitidis strain A antigen encoded by ORF22.
XX
KW Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
treatment; Neisseria infection; meningitis; septicemia; gonorrhea.
XX
OS Neisseria meningitidis.
XX
PN WO9924578-A2.
XX
PD 20-MAY-1999.
XX
PF 09-OCT-1998; 98WO-IB01665.
XX
PR 01-SEP-1998; 98GB-0019016.
PR 06-NOV-1997; 97GB-0023516.
PR 14-NOV-1997; 97GB-0024190.
PR 18-NOV-1997; 97GB-0024386.
PR 27-NOV-1997; 97GB-0025158.
PR 10-DEC-1997; 97GB-0026147.
PR 14-JAN-1998; 98GB-0000759.
XX
PA (CHIR-) CHIRON SPA.
XX
PI Grandi G, Masignani V, Pizza M, Rappuoli R, Scarlato V;

```

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XX
DR WPI: 1999-327407/27.
DR N-PSDB; AA212027.
XX
PT Proteins from Neisseria meningitidis and N. gonorrhoeae useful for
diagnosis, treatment and prevention of infection
XX
PS Claim 4; Page 123; 524pp; English.
XX
CC Amino acid sequences AAV38499-Y38944 represent Neisseria meningitidis
and N. gonorrhoeae antigenic proteins. They are encoded by open
CC reading frames (ORFs) AA211972-212358. The antigenic proteins,
CC their fragments, their nucleic acids and antibodies are used for
CC diagnosis, prevention (as vaccines) or treatment of Neisseria
CC infections, such as meningitis, septicemia and gonorrhea. Both
CC organisms are closely related. Fragments of the nucleic acids
CC are useful as hybridisation probes and antisense reagents.
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XX
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Quality: 2177.00 Length: 447
Ratio: 4.982 Gaps: 0
Percent Similarity: 97.763 Percent Identity: 94.855
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17 uGlnValIleIleTyraSpGlyProValIleThrGluValAlaLeuLeuGlyG 34
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101 AAGAAATATGCCGTATGCCGCCCTCGATGAAGTCAAGGAAGCCATGCC 150
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34 lnglnturyrAlaGlymrGlymrGlymrGlymrGlymrGlymrGlymrGlymr 50
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151 GTCAAAAAAGGCCAAGTCTGTTTGAAGACAAAAGAAATCCGCGGTGCT 200
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51 ValIlelyslsglylGlnValleuPheGluAspIlysls***ProGlymrValVa 67
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201 GTTTACTGCGCGGCTTACAGGCAAAATGCCCGCATTCACCGTGGCGAAA 250
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67 lPheThrAlaProValSerGlyLysIleAlaIleHisArgGlyGluL 84
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
251 AGGCGCTACTCTCAGTGCATGCGATGCGGTGGAAGGCAAGCAAGCAATC 300
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84 ysaGlyValleuGlnSerValIleAlaValaGluGlymrGlymrGlymrGlymr 100
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301 GAGTTGAACGCTACGACCTGGAAGCGCTGGCAAACTTAAGCGGCGAGA 350
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101 GluPheGluArglyrAlaProgluAlaLeuAlaAsnleuSerGly**G1 117
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351 AGTGGCGCGCAACCTGATCCAAATCGGGTTTGTGAGCTGCGTGGCACCC 400
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117 u*****AsnleuIleGlnSerGlyLeuTrpThrAlaLeuArg***A 134
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401 GTTCGTTGAGCAAAATCTGCGCGTGAAGCGGAGCGGCTGGCAATCTTC 450
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134 rglProPheSerlyslleProAlaValaAspAlaGlnProPheAlaIlePhe 150
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
451 GTCAATGCGATGAGACCAATCCGCTGCTGCGGACCCATCGCATATAT 500
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151 ValAsnAlaMetaspThrAsnProleuAlaIlaAspProValValIle 167
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
501 CAAAGAGCGCGCGAGGATTTCAAACGCGGCGCTGTTGGATTAGCCGCTT 550
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[illegible]

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251 AGCGGACTTCACTGACGTCGATTGCGCTTGAAGCAGCAGCAATC 300
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84 ysargValleuGlnSerValValleAlaValGlu***AsnAspGluIle 100
301 GAGTTTGACGCTACGACCTGAAAGCGCTGGCAAACTTAACGGCGAGA 350
      ||||||| ||||||| ||||||| ||||||| ||||||| |||||||
101 GIupheGluArgTyrAlaProGluAlaLeuAlaAsnLeuSerGlyGluG 117
351 AGTGGCGCGCAACGATCCATCCGCTTGTGACCTGCGCTGGCGCACCC 400
      ||||||| ||||||| ||||||| ||||||| ||||||| |||||||
117 uValArGArGAsnLeuIleGlnSerGlyLeuTrpThrAlaLeuArGThrA 134
401 GTCCGTTGACGAAATTCCTGCGCTGATGCCGAGCGCGCTTGGCATCTT 450
      ||||||| ||||||| ||||||| ||||||| ||||||| |||||||
134 rGProPheSerLysIleProAlaValAspAlaGluProPheAlaIlePhe 150
451 GTCAATGCGATGACACCATCCG 474
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151 ValAsnAlaMetAspThrAsnPro 158

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seq_documentation_block:

ID AAV34439 standard; Protein; 451 AA.

AAV34439;

25-AUG-1999 (first entry)

Porphyromonas gingivalis protein PGI.

Porphyromonas gingivalis; Pg; periodontal disease; gingivitis;

vacaine; antigenic.

Porphyromonas gingivalis.

MO929870-A1.

17-JUN-1999.

10-DEC-1998; 98WO-AU01023.

04-AUG-1998; 98AU-0005028.

10-DEC-1997; 97AU-0000839.

31-DEC-1997; 97AU-0001182.

30-JAN-1998; 98AU-0001546.

10-MAR-1998; 98AU-0002264.

09-APR-1998; 98AU-0002911.

23-APR-1998; 98AU-0003128.

05-MAY-1998; 98AU-0003338.

22-JUL-1998; 98AU-0003654.

29-JUL-1998; 98AU-0004917.

(CSLC-) CSL LTD.

Agilus CT, Barr IG, Hocking DM, Margetts MB, Patterson MA;

Ross BC, Rothel LJ, Webb EA;

WPI, 1999-385613/32.

N-PSDB; AAV31657.

Antigenic Porphyromonas gingivalis peptides for preventing

gingivitis

Claim 1; Page 417-418; 588pp; English.

CC be used to detect Porphyromonas gingivalis in standard hybridisation

CC assays. Porphyromonas gingivalis is involved in periodontal disease

CC especially gingivitis.

XX SQ Sequence 451 AA;

alignment_scores: Quality: 663.00 Length: 452
Ratio: 2.225 Gaps: 7
Percent Similarity: 65.929 Percent Identity: 34.735

alignment_block:

us-09-303-518d-125 x AAV34439

Align seg 1/1 to: AAV34439 from: 1 to: 451

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4 ValIleLysThrLysLysGlyLeuAlaLeuAsnLeuLysGlyProIle 20
51 GCAGCGCGTTTACGACGCGCGCCATTAACGAAATC...GGTGGCTTG 97
      ::|||:: |||||:: |||:: |||:: |||::|||
20 uProGluMetLeuAlaGluProAlaGlnSerProThrTyrAlaValAlp 37
98 GCGAGATATATGCCGCTATGCCGCCCTCGATGAAAGTCAAGAGCGAT 147
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37 romspasppheGluGlyValIleProLysValThrAlaTrpGlyasp 53
148 GCGGTCAAAAAGGCCCAAGTGTGTTTGAAGACAAAAAGATCCGGCGT 197
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54 LysValAlaGlaGlySerAlaLeuMetHisLysAlaTyrProGluMe 70
198 GGTGTTTACTGCGCGCGCTTCAAGCAAAATCCCGGATTAACCGTGCG 247
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70 LysPheThrSerProValSerGlyGlyValIleAlaValAsnArgGlyA 87
248 AAAAGCGGCTACTTCAGTCGATGATGCCGCTTGAAGCAAGCAAGCA 297
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87 LysArgLysValLeuSerIleGluValLysProspLysLeuAsnGlu 103
298 ATCGAG...TTTGAAAGCTACGACCTGAAAGCGCTGCAAACTTAAGCG 344
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104 TyrGluSerPheProValGlyAspProSerAla.....LeuSerAl 117
345 CGAAGAGTGGCGCGGCAACCTGATCCATCCGCTTGGAGCTGGCGTGC 394
      :|||::|||::|||::|||::|||:: |||::|||::
117 agluGlnIleLysGluLeuLeuSerSerGlyMetTrpGlyPheIle 134
395 GCACCGCGCTGTCAGCAAAATTCCTGCGCGTGGATGCCGAGCGCTTCC 444
      : |||||::|||:: |||:: |||:: |||||:: |||::
134 yseGlnArgProTyrAspIleValAlaThrProAspIleAlaProArgasp 150
445 ATCTTCGTAATGCGATGACCAATCCGCTGCTGCGCAGCCCTTAACG 494
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151 IleTyrIleThrAlaAsnPheThrAlaProLeuAlaIleProAspPhe 167
495 CATTTCAAAAGAACCGCGGAGATTCAACGCGCGCTGTTGATTGA 544
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167 eIleValArgGlyGluGluArgAlaLeuGlnThrAlaIleAspAlaLeu 184
545 GCGGTTTGACGACGCAAAATTCATGTTTGAAGCAGCTGGCGCAAC 594
      ::|||:: |||||:: |||||:: |||||:: |||||
184 LysLeuThrThrGlyLysValTyrValGlyLeuLysProIleLysSer 200
595 GTCCGCTGAAATGCTGCCAATCGAACATCGAACATCGAATTCGCGCGCC 644
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201 LeuGlyLeuHisAsnAlaGluIleValGluValHis.....GlyPr 214
645 GCATCTGCGCGTTTGATGGAGCGACCATTTATTTCAATCGACGCGTGC 694
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214 ohspProAlaGlyAsnValGlyValLeuIleAsnHisThrLysProIleA 231

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90 lalysalglvsvalleuserillegluvalylsproaspolyleuasglu 106
298 ATTCAG...TTTGACGCTACGACCTGAAGCGCTGGCAACTTACGG 344
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107 TyrGluserPheProValGlyAspProSerAla.....LeuSerAl 120.
345 CGAAGAAATGGCGCGCAACCTGATCAATCCGTTTGTGACCTGCTGC 394
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120 agluGlnIleLysgluleuLeuSerSerGlyMetTrpGlyPheIle 137
395 GCACCCGCTTCGCAAAATTCCTGCCGCTGCCGAGCGCTGGCC 444
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137 ysglnaGpTrpTraspIleValAlaThrProaspIleAlaProasp 153
445 ATCTTCGTCATGCGATGACGACCAATCCGTGGCTGCCGACCTACGGT 494
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154 lIeTyrIleThrAlaAsnPheThrAlaProleuAlaProaspPheasp 170
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495 CATTAATCAAGAGAGCGCGCGAGATTTCAAACGCGCTGTGATTGA 544
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170 eIleValaArgIlygluIaArgAlaLeuGlnThrAlaIleaspAla 187
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187 lalysleuThrThrGlyLysValIlyValGlyLeuLysProGlySer 203
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204 LeuGlyLeuHisAsnAlaGluIleValGluValHis.....GlyPr 217
645 GCATCTCCCGCTTGAAGTGGCAGCGACATTCATTCGACGCGCGCG 694
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217 OhIsProAlaGlyAsnValGlyValIleuLisHisThrLysProIle 234
695 GCGCGAATAAACCCTGTGACCATTCATTCAGATGATTAACCAT 744
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234 snArgGlyGlnThrValIThrPheLysAlaThrAspIleuIleVal 250
745 GCGCGCTTGTTCACACAGCGCGCTGTAACGCGCGCTGATTCCT 794
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251 GlyArgPheLeuThrGlyLysAlaAspPheThrArgMetIleAla 267
795 AGTGCTTCTCAAGTCAACAAACGCGCTCTGCTACCTTTGGTG 844
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267 ThrGlySerAspAlaAlaHisGlyTyrValArgIleMetProGly 284
845 CGAAGTATCGCAATTAAGTGGCGAATGTT.....GACACAGAC 888
||||| ... ||||| ... |||||
284 yAsnValPheAlaSerPheProGlyArgLeuThrIleLysGluSer 300
889 AACGCGTATTCGCGTTCGATTCGATTCGACGCGGATTCACAGGCG 938
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301 GluArgValIleAspGlyAsnValIleThrGlyLysLysLeuGly 317
939 GCACGATTAATTTGGACGCTACCAATCAATCAATTCGTTATTCGAAG 988
||||| ... ||||| ... |||||
317 sGlnProPheLeuSerAlaArgCysAspGlnIleThrValIlePro 334
989 GCGCGAGC...AAAGAGCTGTTCGCTGGTTCGCGCGCGAGCGGCA 1035
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334 lYAspAspValAspGluLeuPheGlyTrpAlaAlaProArgLeuasp 350
1036 TACTCCATCAGCGCTACAACTCGCGCATTTCTCG...AAACAAACT 1082
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351 TyrSerMetSerArgAlaTyrPheSerTrpLeuGlnGlyAsnIly 367
1083 CTTCAGTTCACACAGCGCTACAGCGCGCGCGCGCGCGCGCGCGGA 1132
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367 uTyrValIleuAspAlaArgIleGlyGlyGlnArgIleuAlaMet 384
1133 TTGGTACTACGAGCGCGTATGCTTGGATATCTGCCACCTGCTT 1182
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384 eRAsnGluTyrAspArgValPheProMetAspIleTyrProGluTyrLeu 400
1183 TTGGCGGATTTATGCTGGCATACCAACAGCGCGCGCATTTGGCTG 1232
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401 LeuLysAlaIleIleAlaPheAspIleAspLysMetGluAspLeuGly 417
1233 CTTCGAATTTGGACGAGAACCTCGCTTGTGACGCTTCGTCGCCG 1282
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417 eTyrGluValAlaIleProGluAspPheAlaThrCysGluPheValAsp 434
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AC AA75272;
XX
DT 21-MAR-2000 (first entry)
XX
DE Neisseria meningitidis ORF 628 protein sequence SEQ ID NO:2018.
XX
KW Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
antigenic; diagnosis; immunogenic; infection; meningitis; septicaemia;
antibacterial; gene therapy.
XX
OS Neisseria meningitidis.
XX
PN W09957280-A2.
XX
PD 11-NOV-1999.
XX
PF 30-APR-1999; 99WU-US09346.
XX
PR 01-MAY-1998; 98US-0083758.
PR 31-JUL-1998; 98US-0094869.
PR 02-SEP-1998; 98US-0098994.
PR 02-SEP-1998; 98US-0098062.
PR 09-OCT-1998; 98US-0103749.
PR 09-OCT-1998; 98US-0103794.
PR 09-OCT-1998; 98US-0103796.
PR 25-FEB-1999; 99US-0121528.
XX
PA (CHIR ) CHIRON CORP.
PA (GENO-) INST GENOMIC RES.
XX
PI Fraser C, Galeotti C, Grandi G, Hickey E, Masignani V, Mora M,
PI Petersen J, Pizza M, Rappuoli R, Ratli G, Scalato E, Scarselli M,
PI Tettelin H, Venter JC;
XX
DR WPI: 2000-062150/05.
DR N-PSDB: AA254034.
XX
PT Novel Neisserial polypeptides predicted to be useful antigens for
PT vaccines and diagnostics
XX
PS Claim 2; Page 1003; 1453pp; English.
XX
CC AA253015 to AA254536, AA254577 to AA254615, and AA774253 to AA775941
CC represent novel Neisseria meningitidis and N. gonorrhoeae polynucleotides
CC and polypeptides. AA254537 to AA254576 and AA254616 to AA25473 represent
CC PCR primers used in the exemplification of the present invention. The
CC polypeptides, the polynucleotides, antibodies and compositions of
CC the invention can be used as vaccines, as diagnostic reagents, and as
CC immunogenic compositions. The polypeptides can be used in the
CC manufacture of medicaments for treating or preventing infection due to
```

CC Neisserial bacteria (e.g. meningitis and septicemia), to detect the
 CC presence of Neisseria bacteria, or to raise antibodies. They may also
 CC be used to screen for agonists or antagonists, which may themselves
 CC have use as antibacterial agents. The polynucleotides of the invention
 CC may also be used in gene therapy protocols.

XX Sequence 120 AA;

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 Ratio: 5.000 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-303-518D-125/rev x AAY75272 ..

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624 TTCGATGTTGGCAGCATTTTCAGACGCGACGCTCGCCGACCTGCTTAC 575
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17 IserMetLeuAlaAlaPheSerAspGlyThrSerAlaProAlaLeuG 34
574 AAACATGATTTTGGCTTGGTCAAAACGGCTCAATACCAACAGCCGCGT 525
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seq_documentation_block:

ID AAY75273 standard: Protein: 120 AA.

XX AAY75273;
 XX
 XX 21-MAR-2000 (first entry)
 XX
 DE Neisseria meningitidis ORF 628 protein sequence SEQ ID NO:2020.
 XX
 XX Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
 KW antigenic; diagnosis; immunogenic; infection; meningitis; septicemia;
 KW antibacterial; gene therapy.
 XX
 OS Neisseria meningitidis.
 XX
 PN WO957280-A2.
 XX
 PD 11-NOV-1999.
 XX
 PF 30-APR-1999; 99WO-US09346.

XX
 PR 01-MAY-1998; 98US-0083758.
 PR 31-JUL-1998; 98US-0094869.
 PR 02-SEP-1998; 98US-0098994.
 PR 02-SEP-1998; 98US-0099062.
 PR 09-OCT-1998; 98US-0103749.
 PR 09-OCT-1998; 98US-0103794.
 PR 09-OCT-1998; 98US-0103796.
 PR 25-FEB-1999; 99US-0121520.
 XX

PA (CHIR) CHIRON CORP.
 PA (GENO-) INST GENOMIC RES.

PI Fraser C, Galeotti C, Grandi G, Hickey E, Masignani V, Mora M,
 PI Petersen J, Pizzia M, Rappuoli R, Ratti G, Scalato E, Scarselli M,
 PI Tettelin H, Venter JC;

DR WPI; 2000-062150/05.
 DR N-PSDB; AA254035.

PT Novel Neisserial polypeptides predicted to be useful antigens for
 PT vaccines and diagnostics

PS Claim 2; Page 1004; 1453pp; English.

XX
 XX AA253015 to AA254536, AA254577 to AA254615, and AAY74253 to AAY75941
 CC represent novel Neisseria meningitis and N. gonorrhoeae polynucleotides
 CC and polypeptides. AA255537 to AA254576 and AA254616 to AA255473 represent
 CC PCR primers used in the exemplification of the present invention. The
 CC polypeptides, the polynucleotides, antibodies and compositions of
 CC the invention can be used as vaccines, as diagnostic reagents, and as
 CC immunogenic compositions. The polypeptides can be used in the
 CC manufacture of medicaments for treating or preventing infection due to
 CC Neisserial bacteria (e.g. meningitis and septicemia), to detect the
 CC presence of Neisseria bacteria, or to raise antibodies. They may also
 CC be used to screen for agonists or antagonists, which may themselves
 CC have use as antibacterial agents. The polynucleotides of the invention
 CC may also be used in gene therapy protocols.

XX Sequence 120 AA;

alignment_scores:
 Quality: 574.00 Length: 120
 Ratio: 4.824 Gaps: 0
 Percent Similarity: 99.167 Percent Identity: 95.000

alignment_block:

US-09-303-518D-125/rev x AAY75273 ..

Align seg 1/1 to: AAY75273 from: 1 to: 120

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674 ATGTCGCGCCACTCAAAACCGGAGATGCGGCGCCGGAATTCATGTGT 625
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1 MetCysValProLeuLysProAlaGlyCysGlyProPheSerCysVal 17
624 TTCGATGTTGGCAGCATTTTCAGACGCGACGCTCGCCGACCTGCTTAC 575
|||||
17 IserMetLeuAlaAlaPheSerAspGlyThrSerAlaProAlaLeuH 34
574 AAACATGATTTTGGCTTGGTCAAAACGGCTCAATACCAACAGCCGCGT 525
|||||
34 IstThrPileuLeuArgSerValLysArgLeuAsnThrSerLysProArg 50
524 TTGAATCTCTGGCGGCTCTTTGATTAATGACCGTAGGGTCGCGACG 475
|||||
51 LeuYssSerSerAlaAlaSerLeuIleMetThrValGlySerAlaAla 67
474 CGGATGTCGTCATCGCATTCAGACGATGCGCAAGCGCTCGGATCGA 425
|||||
67 rGlyLeuValSerIleAlaLeuThrLysMetAlaAsnGlySerAlaSer 84
424 CGGACGAAATTTGCTGTAACGAGCGGTCGCGACGCGACGATCCACAACG 375

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|||||
84 hrhlaaglylleuleuasncllyargvalargserhlaValhlslypro 100
374 GATTGGATCAGGTTGCGGCGCAGCTTCTTCGCCGTTAAGTTGGCAGCGC 325
|||||
101 AsprtlleargleuArgThrserSerProleuLysPhehlaAsnal 117
324 TTCAGGTGCG 315
|||||
117 aserGlyAla 120

seq_name: /SIDS1/gcgdata/geneseq/geneseqp-emb1/AA2000.DAT:AAV75271
seq_documentation_block:
ID AAV75271 standard; Protein; 119 AA.
XX
AC AAV75271;
XX
DT 21-MAR-2000 (first entry)
XX
DE Neisseria gonorrhoeae ORF 628 protein sequence SEQ ID NO:2016.
XX
KW Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
KW antigenic; diagnosis; immunogenic; infection; meningitis; septicemia;
KW antibacterial; gene therapy.
XX
OS Neisseria gonorrhoeae.
XX
PN WO957280-A2.
XX
PD 11-NOV-1999.
XX
PF 30-APR-1999; 99WO-US09346.
XX
PR 01-MAY-1998; 98US-0083758.
PR 31-JUL-1998; 98US-0094869.
PR 02-SEP-1998; 98US-0098994.
PR 02-SEP-1998; 98US-0099062.
PR 09-OCT-1998; 98US-0103749.
PR 09-OCT-1998; 98US-0103794.
PR 09-OCT-1998; 98US-0103796.
PR 25-FEB-1999; 99US-0121528.
XX
XX (CHIR ) CHIRON CORP.
XX PA (GENO-) INST GENOMIC RES.
XX
PI Fraser C, Galeotti C, Grandi G, Hickey E, Maignani V, Mora M;
PI Petersen J, Piza M, Rappuoli R, Ratti G, Scalato E, Scarselli M;
PI Tettelin H, Venter JC;
XX
XX WPI: 2000-062150/05.
XX DR N-PSDB; AA254033.
XX
XX Novel Neisserial polypeptides predicted to be useful antigens for
XX PT vaccines and diagnostics
XX PS
XX Claim 2; Page 1003; 1453pp; English.
XX
XX AA253015 to AA254536, AA254577 to AA254615, and AAV74253 to AAV75941
XX represent novel Neisseria meningitidis and N. gonorrhoeae polynucleotides
XX and polypeptides. AA254537 to AA254576 and AA254616 to AA25473 represent
XX PCR primers used in the exemplification of the present invention. The
XX polypeptides, the polynucleotides, antibodies and compositions of
XX the invention can be used as vaccines, as diagnostic reagents, and as
XX immunogen compositions. The polypeptides can be used in the
XX manufacture of medicaments for treating or preventing infection due to
XX Neisserial bacteria (e.g. meningitis and septicemia), to detect the
XX presence of Neisseria bacteria, or to raise antibodies. They may also
XX be used to screen for agonists or antagonists, which may themselves
XX have use as antibacterial agents. The polynucleotides of the invention
XX may also be used in gene therapy protocols.
XX
XX Sequence 119 AA:

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alignment_scores:
Quality: 538.50 Length: 119
Ratio: 4.642 Gaps: 1
Percent Similarity: 97.479 Percent Identity: 93.277

alignment_block:
US-09-303-518D-125/rev x AAV75271 ..

Align seg 1/1 to: AAV75271 from: 1 to: 119

674 ATGTGGTGGCTCAACCGCAGGATGCGGCGCGCCGAATTCATGTGT 625
|||||
1 MetCysValProleuLysProAlaGlyCysGlyProProAsnSerCysVal 17
624 TTCGATGTGGCAGCATTTTTCAGACGCGCTCTGCCCAAGCTTTCAC 575
|||||
17 lserlleuAlaAlaPheSerAspGlyThrSerAlaProAlaAlaLeuH 34
574 AACATCGATTTGCGTTCGTCGAACGGCTCAATACCAAGCGCGGT 525
|||||
34 lstrtrpilleuValSerValArgleuAsnThrSnrArgProAlaG 50
524 TTGAATCTCTGCGCGCTTCTTGATATGACCGTAGGGTCGCGACCGAG 475
|||||
51 leuLysSerSerAlaAlaSerleuMetMetThrValGlySerAlaAla 67
474 CGGATGGTGTCATCGCATTCGACGAAGATGGCGAAGCGCTCGCATCA 425
|||||
67 rglyleuValSerlleAlaLeuThrlysmetAlaAsnGlySerAlaSerT 84
424 CGGCGAGATTTGCTGACGCGAGCGGCTGCGCAGCGAGCCCAACCG 375
|||||
84 hrhlaaglylleuleuasncllyargvalargserhlaValhlslypro 100
374 GATTGGATCAGGTTGCGGCGCAGCTTCTTCGCCGTTAAGTTGCCAGCGC 325
|||||
101 Asp...lleArgleuArgArgThrPheSerleuLeuAsnPhelaAlaSerAl 116
324 TTCAGGT 318
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116 aserGly 118

seq_name: /SIDS1/gcgdata/geneseq/geneseqp-emb1/AA2000.DAT:AAV82082
seq_documentation_block:
ID AAV82082 standard; Protein; 467 AA.
XX
AC AAV82082;
XX
XX 01-JUN-2000 (first entry)
XX
XX Chlamydia pneumoniae antigen CPN100605 protein SEQ ID NO:2.
XX DE Chlamydia pneumoniae; antigen; CPN100605 protein; immunisation;
XX KW vaccine; infection; antibacterial; antiinflammatory; bronchitis;
XX KW community acquired pneumoniae; upper respiratory tract infection;
XX sinusitis.
XX
XX Chlamydia pneumoniae.
XX OS
XX PN WO200006742-A2.
XX
XX 10-FEB-2000.
XX
XX 27-JUL-1999; 99WO-IB01331.
XX PF
XX 27-JUL-1998; 98US-0094195.
XX PR
XX 26-JUL-1999; 99US-0361443.
XX
XX (CONN-) CONNAUGHT LAB LTD.
XX
XX

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PI Mardin AD, Oomen RP;
XX WPI: 2000-205466/18.
DR N-PSDB; AAZ95378.
XX
PT Chlamydia pneumoniae antigens used for immunization and protection
PI against Chlamydia diseases -
XX
PS Claim 6; Fig 1; 48bp; English.

CC The present sequence represents the Chlamydia pneumoniae antigen
CC CPN100605 protein. The CPN100605 protein has antibacterial and
CC antiinflammatory activities. The Chlamydia pneumoniae CPN100605
CC polynucleotide and protein can be used in vaccination methods for
CC preventing and treating Chlamydia infection (e.g. infections caused by
CC C. trachomatis, C. psittaci, C. pneumoniae or C. pecorum). The
CC polynucleotide can be used to produce the protein recombinantly, in the
CC construction of vaccine vectors, as a vaccine agent, and in the
CC construction of an attenuated Chlamydia strain. The protein are also be
CC useful as a vaccine agent, and for the preparation of medicaments for
CC treating or preventing Chlamydia infection, e.g. community acquired
CC pneumonia, and upper respiratory tract infections such as bronchitis and
CC sinusitis.

SO Sequence 467 AA;

alignment_scores:
Quality: 464.50 Length: 464
Ratio: 1.585 Gaps: 15
Percent Similarity: 63.147 Percent Identity: 30.172

alignment_block:
US-09-303-518D-125 x AAY82082 ..

Align seg 1/1 to: AAY82082 from: 1 to: 467

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3 Iletlrvalasnaarglyleuasprenleuenglyserprolysgl 19
|||||.....|.....|.....|.....|.....|.....|.....|
51 GCAAGCCGTTTACGAC.....GCGCCGCGCATACCGAAGTCGCGTTC 94
|||||.....|.....|.....|.....|.....|.....|.....|
19 userlypherlyrasnlyslleasprrogluphvalserlleaspreu. 35
|||||.....|.....|.....|.....|.....|.....|.....|
95 TTGGCGAAGATATGCCGCTATGCGCCCTCGATGAAGTCAAGAAAGC 144
|||||.....|.....|.....|.....|.....|.....|.....|
36 .....|.....|.....|.....|.....|.....|.....|
145 GATGCCGTCAAAAAGCCCAAGTGTCTTTAAGACAAAAAAGATCCGGG 194
|||||.....|.....|.....|.....|.....|.....|.....|
51 Aspalavalcysserglyalaproillealaglutyryshisphreproas 67
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195 CGTGTGTTTACTGCGCGGCTTCAGCGAAATGCGCGCATTCACCGTG 244
|||||.....|.....|.....|.....|.....|.....|.....|
67 nthrtyrlethrserhisvalserglyvalthralallethrargg 84
|||||.....|.....|.....|.....|.....|.....|.....|
245 GCGAAACCGCGTACTTCAGTACGTGTAT...GCCGTGAAGGCAAC 291
|||||.....|.....|.....|.....|.....|.....|.....|
84 lyaslylarserserleuaspvalillellyslsthrproglpro 100
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292 GACGAAATCGAGTTGAACGCTACGACCTGAACGCTGGCAACTTAAG 341
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101 Thrsertlrgluty.....Thrtyraspreuenglnthrlleuse 113
|||||.....|.....|.....|.....|.....|.....|.....|
342 CGGCGAAGAAATGCGCGCAACTGATCACTCGGTTTGGAGTCGCG 391
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113 rargseraspreuaserglyleuhsnglyleuphealeu 130
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392 TGCGCACCGCTCGTTGACGAAATTCCTCGCGTGCAGTCCGAG...CCG 438
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130 lelysglnargprophasp...lleproalleleprothrlnthrlpro 145
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439 TTGCGCATTTGTCATGCGATGACACCAATCCGCTGCGCGACCC 488
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146 Argasryvalrheilleasnlleasprasnprrothrlthrlproser 162
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489 TACGGTCATTATC.....AAGAAGCCCGCGAGATP 520
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162 oglylshisleuallaleupheserarglgluglylphetyalr 179
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521 TCAACCGCGCGCTGTGTATGACCGCTTGACCCGCAACCAATTCAT 570
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179 hevalvalglyvalarglallealalalyseuheglyleuargronis 195
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571 GTTGTGTAAGCAGCTGCGCGACAGCTGCGTGAATGCTGCAACAT 620
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196 llevalrheargasrarglyleuphrleprothrlleuylsthrll 212
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621 C...GAACACATGATTCGCGCGCGCGCATTCGCGGTTTGCATGCA 667
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212 ealnlshleuhsThrvalserglyprophroserglyserproserl 229
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668 CGSACATTCATTCATCGACCGCGTGGCGCGCAATAA...ACCGTGG 714
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229 lehlstlehlsserValalaproillehrasnlyslnglylvalrhe 245
|||||.....|.....|.....|.....|.....|.....|.....|
715 ACCATCAATATGACATGTAATTCATTCATTCGCGGTTTGCATGCA 764
|||||.....|.....|.....|.....|.....|.....|.....|
246 Thrleuserhehlinsryallethrllleglynlshleupheleuylsgl 262
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765 CGCTGTGACACCGACGCGCGTGAATGCCCTAGCGTGTTCATGCAACA 814
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262 yarglileuhsnglylvalrhlaleuallaglylthrlaleuyls 279
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815 AACCG.....CGCGTTCGCGTACCGCTTTGGGTGCGCAATGTCGCA 858
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279 erserleuargarglyvallethrthrllysllylaserpheser 295
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859 ...ATTATCGCGCGCAATTCGTTGACACACCAACCGCGTATTCGCG 905
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296 leuileasleuasnprlleseraprasnprthr...leuilesergl 311
|||||.....|.....|.....|.....|.....|.....|.....|
906 TTGCGTATTCGACGCGCGCATTCACACAGCGCGCGCAT...TATTTGG 952
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311 yasprroleuThrnglylrgleucylslslnglgluphroheleug 328
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953 GACGTACCAACATGATTCGTTATGCAAGAGCGCGCAAGAG 1002
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328 lyrheargasprhlsserllesevalleuhsasprrothrllysrarglu 344
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1003 CTGTTGCGCTGGGTTGCGCGCGACCGGCAAAATCTCCATGACCGCTAC 1052
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345 leupheserphleuarglileglyrheasnllysprrrothrlthrllyst 361
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1053 AACCCTCGCGCATTTCTGAAAAACCAACTCTTCAAGTTCAAC.....A 1096
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361 rtyrleuaserglyrhephelelysllyslsargthrtlythrhasnproaspr 378
|||||.....|.....|.....|.....|.....|.....|.....|
1097 CAGCCGTCAACGCGCGCGACCGCGCATGTCGCGATTCGCTACTACGAG 1146
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378 hrasnleuhsnglyluphrargprolleleasprthrasplelytrsp 394
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1147 CGCGTATGCGCTGGATATCTGCGCGACCGCTGCTTTGGCGGATTAAT 1196
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395 lyvalmetrmetelargllepvalvalproleuilelysalavall 411
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1197 GCGCGCGATTCACGACGCGCGCGCATTCGCTGCTGCAATTCGACG 1246
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411 ethrllysasprheasprleuallasnnglyleuhsnglylvalcyg 428
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1247 AAGAAGACCTCGCTTTGACAGCTTCGTCGCGCGCAATACGAAATAC 1296
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428 lygluasprhealaleuoprothrlleuileasprroserlysthrclmet 444
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1297 GCGCGCGCTGTTGCGCAAGTGTGCAAAACCATTCGAAAGGAA 1338
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445 LeuThrIleValIysGluSerLeuIleGluTyrAlaIysGlu 458

seq_name: /STDS1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AA1935375

seq_documentation_block:

ID AA1935375 standard; Protein; 469 AA.

AC AA1935375;

DT 13-SEP-1999 (first entry)

DE Chlamydia pneumoniae transmembrane protein sequence.

KW Respiratory disease; pneumonia; bronchitis; heart disease; sarcoidosis;

KW sinusitis; purulent otitis media; erythema nodosum; pharyngitis;

KW vaccine; neutralising epitope.

OS Chlamydia pneumoniae.

PN W09927105-A2.

PD 03-JUN-1999.

PF 20-NOV-1998; 98WO-IB01890.

PR 04-NOV-1998; 98US-0107078.

PR 21-NOV-1997; 97PR-0014673.

PA (GEST) GENSET.

PI Grifffals R;

DR WPI; 1999-357842/30.

PT Genome sequence of Chlamydia pneumoniae

PS Page 1170-1171; Disclosure; 1912pp; English.

XX AA193584-Y35879 represent the proteins encoded by all the open reading

CC frames in the complete genome (see AA191990) of Chlamydia pneumoniae.

CC C. pneumoniae causes respiratory disease such as pneumonia and

CC bronchitis and is thought to be a contributing factor in heart

CC disease, sarcoidosis, sinusitis, purulent otitis media, erythema

CC nodosum or pharyngitis. The polypeptides encoded by the open reading

CC frames of the C. pneumoniae genome (see AA193584-Y35879) can be used in

CC immunogenic compositions as vaccines. Vectors containing C. pneumoniae

CC nucleotide sequences can also be used as immunogenic compositions,

CC especially where the vector directs the expression of a neutralising

CC epitope of C. pneumoniae.

XX

XX

XX

XX

XX

alignment_block:

US-09-303-518D-125 x AA1935375

Align seg 1/1 to: AA1935375 from: 1 to: 469.

alignment_scores:

Quality: 464.50 Length: 464

Ratio: 1.585 Gaps: 15

Percent Similarity: 63.147 Percent Identity: 30.172

4 ATTAATAATCAAAAAAGTCTTAACCTGCCATCGGGCAGACG...GA 50

5 IletHrValAsnArgGlyLeuAspLeuSerLeuGlnGlySerProLysG1 21

51 GCAACCGCTTACGAC...GGCCGGCATTACCGAAGTCGCGTTCG 94

21 UserIlyPheTyrAsnLysIleAspProGluPheValSerIleAspLeu 37

95 TTGGCAGAAATATGCGCGTATGCGCCCTCGATGAAGTCAAGAGAGGC 144

38ArgProPheGlnProLeuSerLeuLysValGluGlnGly 52

145 GATGCCGCTAAAAAGGCCAATGCTGTTTGAAGCAAAAAAGATCCGG 194

53 AspAlaValCysSerGlyAlaProIleAlaGluTyrLysHisPheProAs 69

195 CGTGGTGTACTACGGCCGCTTACGCAAAATCCGGGATTCACCGTG 244

69 nHrTyrIleThrSerHisValSerGlyValValIleAlaIleArgG 86

245 GCGAAAGCGCGTACTTCAGTCAGTCGAT...GCCGTGAAGCAAC 291

86 LysnLysArgSerLeuAspValIleIleLysThrProLysPro 102

292 GACGAATGAGTTTGAACGCTACGACCTGAGCGCTGGCAACTTAA 341

103 ThrSerThrLysLysLysLysLysLysLysLysLysLysLysLys 115

342 CGCGCAAGAGTGGCGCCGACCTGATCCATCCGCTTGTGACTGCGC 391

115 rArgSerAspLeuSerGluIlePheLysGluAsnLysLeuPheAlaLeu 132

132 LysGlnArgProPheAsp...IleProAlaIleProThrGlnThrPro 147

439 TTGCGCATCTGCTCATGCGATGAGACCAATCCGCTGCGCGACCC 488

148 ArgAspValPheIleAsnLeuAlaAspAsnArgProPheThrPro 164

489 TACGTCATTTATC.....AAGACGCCCGCGAGAT 520

164 OGILYSHISLeuAlaLeuPheSerSerArgGluGluGlyPheTyrVal 181

521 TCAACGCGCGCTGTTGATGAGCGGTTGACGCAACCAAAATCCAT 570

181 heValValGlyValArgAlaIleAlaLysLeuPheGlyLeuArgProHis 197

571 GTTTGTAAGCAGCTGGCCAGACGTCGCTGAAAATGCTGCCAATC 620

198 IleValPheArgAspArgLeuThrLeuProThrGlnGluLeuLysThr 214

621 C...GAACACATGAATTCGGCGCGCCGATCCGCTGAGTGAAGCA 667

214 eAlaHisLysLeuHisThrValSerGlyProPheProSerGlySerProSer 231

668 CGCACATTCATTATCGAAGCGCGTGGCGCAATAA...ACCGTGG 714

231 leHisIleHisSerValAlaProIleThrAsnGluLysGluValAlaPhe 247

715 ACCATCAATATCAAGATGAATGATTAACATTCGCGCTTGTGCAACAG 764

248 ThrLeuSerPheGlnAspValLeuThrIleGlnLysLeuPheLysG1 264

765 CCGCTGACACCGCAGCGGTGATTCCTAGGTGGTTTCAAGTCAACA 814

264 YArgIleLeuHisGluGlnValThrAlaLeuAlaGlyThrAlaLeuLys 281

815 AACCG.....CGCTGTGCGTACCGTTTGGGTGCGAAGTATCGCA 858

281 eSerLeuArgArgTyrValIleThrThrLysGlyLysSerPheSer 297

859 ...ATTACTGCGCGCATTCGTTGACACAGCAACCGGTGATTCGCG 905

298 LeuIleAsnLeuAsnLysPheSerAspAsnAspThr...LeuIleSer 313

906 TTGCGTATTAAGCGCGCATTCACACAGCGCGCAGAT...TATTTGG 952

313 YAspProLeuThrGlyArgLeuLysLysLysGluGluGluProPheLeuG 330

953 GCGCATCAACCAATCAGATTCGTTATCGAAGAGCGCGCAAGAAAG 1002


```

1  ...  |||  ...  |||||  |||||  ...
212 eAlaHisLeuHisThrIleAspGlyProPheProSerGlySerProSerT 229
668 CGCACATTTCATTCATCGAGCCGGTC...GGCCGGAATTAACCGCTGG 714
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229 hrHisIleHisIleHisIleAlaArgIleArgAsnGluArgAspValValPhe 245
715 ACCATCATATTTCAGATGTAATTCATTCATTCATTCATTCATTCATTC 764
|||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
246 ThrIleSerPheGlnGluValLeuSerIleGlyHisLeuPheLeuLysG 262
765 CCGCTGTAACACGAGCGCGGATTCGCTGATTCGCTGATTCGATTCG 810
|||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
262 yPheValLeuLysGlnGlnIleValAlaLeuAlaGlySerAlaLeuPro 279
811 ..AACAAACCGCGCTCTGCGCTGCGCTGCGCTGCGCTGCGCTGCG 858
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279 roSerGlnArgLysThrLeuLeuThrAlaLysGlyAlaSerPheSerAsp 295
859 ATTACTGGGGCGAATTGGTGTACACAGACAAAC...CGCGGATTCGCG 905
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296 LeuLeuProLysAspIlePheSerSerAspIleThrLeuIleSerG 312
906 TTCGGTATTGAACGCGCGGATTCACACAGCGCGCGACGAT...TATTTGG 952
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312 yAspProLeuThrGlyArgLeuLysLysGlnGlnAsnProCysLeuG 329
953 GACGCTACCAATCAGATTCCTGCTTATTCAGAAAGCGCGACGAAAGAG 1002
|||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
329 yMetArgAspHisThrIleThrLeuLeuProAsnProLysThrArgGlu 345
1003 CTGTTGCGGCTGGGTGGCGCGGACCGGCAATATCTCATCAGCGGAC 1052
|||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
346 SerPheSerPheLeuArgLeuGlyTyrAsnLysLeuThrValThrArg 362
1053 AACCTCGCGCATTCCTGTAACAAACAA...CTCTTCAAGTTCAACA 1096
|||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
362 rTyrLeuSerGlyPhePheLysArgLysArgValPheMetAspMetAsp 379
1097 CAGCGCTACAGCGCGCGACCGCGCCAGCGCGCTGCTGCTGCTGCTG 1146
|||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
379 hrAsnMetHisGlyGluLysArgProIleIleAspAlaGluIleTyrGlu 395
1147 GCGGTGATGCGCTGGATATCTGCGCACCGCTGCTGCGGATTTAT 1196
|||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
396 ArgValSerAlaIleProValProValAlaLeuIleIleLysAlaLeuG 412
1197 CGTGGCGCATTCACGACGCGCGACGCGCATTCGCTGCTGCTGCTG 1246
|||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
412 urThrGlnAsnPheGlnGluAlaLysArgLeuLysLeuLeuGluValAla 429
1247 AAGAAAGACCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1296
|||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
429 roGlnAsnPheAlaLeuProThrPheIleAspProSerLysThrGluMet 445
1297 GGCGCGCTGCTGCGCAAGTCTGCTGCTGCTGCTGCTGCTGCTGCTG 1335
|||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
446 PheSerIleValLysGlnSerLeuLeuArgThrGlnLys 458

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seq_name: /SIDSL/gcdata/geneseq/geneseq-emb1/AA1999.DAT:AAV34467

seq_documentation_block:

ID AAV34467 standard; Protein: 443 AA.

AAV34467;

25-AUG-1999 (first entry)

Porphyromonas gingivalis protein PG12.

Porphyromonas gingivalis; PG; periodontal disease; gingivitis;

vaccine; antigenic.

XX

OS Porphyromonas gingivalis.

XX W09929870-A1.

XX 17-JUN-1999.

XX 10-DEC-1998; 98W0-AU01023.

XX 04-AUG-1998; 98AU-0005028.

XX 10-DEC-1997; 97AU-0000839.

XX 31-DEC-1997; 97AU-0001182.

XX 30-JAN-1998; 98AU-0001546.

XX 10-MAR-1998; 98AU-0002264.

XX 09-APR-1998; 98AU-0002911.

XX 23-APR-1998; 98AU-0003128.

XX 05-MAY-1998; 98AU-0003338.

XX 22-MAY-1998; 98AU-0003654.

XX 29-JUL-1998; 98AU-0004917.

XX (CSLC-) CSL LTD.

XX Agnus CT, Barr IG, Hocking DM, Margetts MB, Patterson MA;

XX Ross BC, Rothel LJ, Webb EA;

XX WPI; 1999-385613/32.

XX N-PSDB; AAX91685.

XX Antigenic Porphyromonas gingivalis peptides for preventing

XX gingivitis

XX Claim 1; Page 445-446; 588pp; English.

XX AAX91536 to AAX91801 encode two hundred and sixty six antigenic

XX Porphyromonas gingivalis (PG) polypeptide sequences given in AAV34318 to

XX AAV34583. AAX91802 to AAX91989 represent PCR primers used in the

XX isolation of the PG polypeptides. The PG polypeptides have antibacterial

XX activity with a vaccine mechanism of action. The PG polypeptides can be

XX used as vaccines especially against Porphyromonas gingivalis. Probes can

XX be used to detect Porphyromonas gingivalis in standard hybridisation

XX assays. Porphyromonas gingivalis is involved in periodontal disease

XX especially gingivitis.

XX Sequence 443 AA;

XX alignment_scores:

XX Quality: 169.50 Length: 483

XX Ratio: 0.712 Gaps: 21

XX Percent Similarity: 49.275 Percent Identity: 20.290

XX alignment_block:

XX US-09-303-518D-125 x AAV34467 ..

Align seg 1/1 to: AAV34467 from: 1 to: 443

37 GCGGCGACGCGGAGCAAGCGTTTACGAGCGCCGCGCATTCACGAAGT 86

19 AlaGlyLysProValGluValLeu.....ProIleProSerGlnVal 32

87 GCGCTGCTGGCGCAAGATATGCGGATGCGCGCTTCATCATCAAGTCA 136

32 lValIleProLeuGlnHisIleGlyAlaProAlaThrAlaThrVal 49

137 AGGAAGGCGATGCGCTCAAAAAGCCAGCGCTTGAAGACAAAAG 186

49 yLysGlyAspGluValValValGlyThrIle..... 60

187 AATCCGGCGCTGTGTTTACTGCGCGCTTCAGCAAAATCGCGCG.. 234

61AlaGlnAlaGlyLysPheValSerAlaAs 70

235 .ATTACCGCTGGC...GAAAGCGGCTACTGAC..... 264

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70 nllhlsrserValSerGlyValLeuLysIleaspasValTYRA 87
265 .....TCAGTCGATGTCGGCTGGAAGGACAC 291
87 spSerSerGlyTyrProLysProAlaValPheIleSerValIleGlyasp 103
292 GACGAATCGAGCTTGAACGCTACGACCGATGAAGCGCTG.....GCAAA 335
::: ||| ::|
104 GIUTPrGIUGLyIleaspSerProAlaIleValLysGIUCysAs 120
336 CTTAGCGGCGGAGAGAGTGGCCGCAACCTGATCCATCCGCTTGTGGA 385
||||| ::|
120 nleuaspAlaLysGIUIleValAlaLysIleSerAlaIleGlyIle...V 136
386 CTGCGCTGCGACCGCTCCGTTTC.....AGCAAAATT 417
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136 aIGlyLeuGIyGIyAlaThrPheProThrHisValLysLeuSerProPro 152
418 CCGCGCGTCGATGCGGACCGCTTGCCCATCTTCGTCATGCGATGACAC 467
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153 ProGIaSnLysAlaGlu.....IleLeuIleIleasnAlaValGIUCy 167
468 CAATCCGCTGGCTGCCGACCTACGACCTATATCAAGAAGACCGCGGAG 517
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167 sGIuProTyrIleuThrSerAspHisValLeuMetLeuGluHisGIyGIUG 184
518 ATTTCAAACGCGCGCTGTGTGATTGAGCCGTTGACCGAAGCAAAATC 567
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184 IUIleMetIleGIyAlaSerIleLeuMet.....LysAlaIle 196
568 CATGTTGTAGGAGCT...GGCGGACAGCTGCCGCTGCTGAAAATGCT.. 612
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197 GIuValasnLysAlaValIleGIyValGIuasnAsnLysAspAlaIle 213
613 .....GCCAACATCGAAACACATG 631
213 eAlaHisLeuThrLysLeuAlaThrAlaTyrProGIyIleGIuValMetP 220
632 AATTCGGCGCGCGCATCCCTGCGGCTTGAGTGGACGACCATTCATTTC 681
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230 rOIeLysValGIuTyrProGIuGIyGIyLysGIuLeuIleaspAla 246
682 ATC.....GAGCGGCTGGCGCGGAATA 704
::: ||| ::|
247 ValIleArgLysGIuValLysSerGIyAlaLeuProIleSerThrGIyAl 263
705 AACCGTGTGACCATCATATATCAAGATGTAATTACCATGGCCGTTTGT 754
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263 aValVal.....GIuasnValGIyThrValPheAlaValI 275
755 TTGCAACAGCGCGCTGTAAC.....ACGAGCGCGCTGATGCCCTTA 795
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275 YrGIuAlaValGIuLysAsnLysProLeuValGIuArgIleValIThrVal 291
796 GGTGGTCTCAAGTCAACAACCGCGCTTGTGCTACGCTTGTGGTGC 845
||||| ::|
292 ThrGIyLysLysLeuSerArgProSerAsnLeuLeuValArgIleGIyTh 308
846 GAAAGATCGCAAAATT.....ACTGCGGCGCAATTGGTTGACACAGACA 889
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308 rProIleAlaIleAlaLeuIleGIuAlaIleGIyGIyLeuProGIuAsnThrG 325
890 ACCGCGGATTCGGGTTGGTATTGAACGGCGCATTCACACAGCGCGC 939
::: ||| ::|
325 LysLysIleIleGIyGIyGIy..... 331
940 CACGATTATTGGAGCGCTACACATCAGATTTCGTTATGAGAGAAG 989
331 ..... 331
990 CCGCAGCAAGAGCTGTGCGCTGGGTTGCGCGACCGCAGCAAAATACT 1039
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332 .....PrometMetGIyArgAlaLeuLeuSerProasp...ValP 344

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1040 CCAFCACGCGTACACACCTTCGCCATTTCCTGAAAAACAACACTTTCAAG 1089
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1090 TTCACACAGCGCGTCAACAGCGCGCGACCGCGCATGCTGCCATTGTGAC 1139
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358 ..GIUGIuAlaValAlaArgLysPrometArgAspCysIleArgCysAlaLy 373
1140 TTACAGACGCGGTGATGTCCTTGGATATCCGCCACACCTGCTTTGCGCG 1189
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373 scYsValGIyValCysPrometGIyLeuasnProAlaPheLeuMetArgA 390
1190 ATTATATCGTGGCGCATACCGACAGCGCGCAG.....GCATTGGGT 1230
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390 sPhrITrLysSerITrPGIuThrAlaGIuLysGIyAsnValValasp 406
1231 TGCTTGGAATTGGACAGAGAGACCTCGCTTGTGCGAGCTGTCGCGCC 1280
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407 CysIleGIuLysGIySer.....CysSerPheThrCysPr 418
1281 GCGCAAAATCGAATACGCGCGCTGTGCGCAAACTGCTGAAACATT 1329
::: ||| ::|
418 oAlaasnArgProLeuLeuAspTyrIleArgGIuAlaLysLysThrVal 434

seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AA134343
seq_documentation_block:
ID AA134343 standard; Protein; 451 AA.
AA134343;
25-AUG-1999 (first entry)
DE Porphyrymonas gingivalis protein Pg122.
KW Porphyrymonas gingivalis; Pg; periodontal disease; gingivitis;
  vaccine; antigenic.
OS Porphyrymonas gingivalis.
  XN W09929870-A1.
  XX 17-JUN-1999.
  XX 10-DEC-1998; 98WO-AU01023.
  XX 04-AUG-1998; 98AU-0005028.
  XX 10-DEC-1997; 97AU-0000839.
  XX 31-DEC-1997; 97AU-0001182.
  XX 30-JAN-1998; 98AU-0001546.
  XX 10-MAR-1998; 98AU-0002264.
  XX 09-APR-1998; 98AU-0002911.
  XX 23-APR-1998; 98AU-0003128.
  XX 05-MAY-1998; 98AU-0003358.
  XX 22-MAY-1998; 98AU-0003654.
  XX 29-JUL-1998; 98AU-0004917.
  XX
  XX (CSLC-) CSL LTD.
  XX
  XX Agius CT, Barr IG, Hocking DM, Margets MB, Patterson MA;
  XX Ross BC, Rothel LJ, Webb EA;
  XX
  XX WPI; 1999-385613/32.
  XX DR N-PDB; AA191561.
  XX
  XX Antigenic Porphyrymonas gingivalis peptides for preventing
  XX gingivitis
  XX
  XX Claim 1; Page 303; 588pp; English.
  XX
  XX AA191536 to AA191801 encode two hundred and sixty six antigenic
  XX Porphyrymonas gingivalis (Pg) polypeptide sequences given in AA134318 to
  CC

```


CC AAY34583. AAY91802 to AAY91989 represent PCR primers used in the isolation of the PG polypeptides. The PG polypeptides have antibacterial activity with a vaccine mechanism of action. The PG polypeptides can be used as vaccines especially against Porphyromonas gingivalis. Probes can be used to detect Porphyromonas gingivalis in standard hybridisation assays. Porphyromonas gingivalis is involved in periodontal disease especially gingivitis.

XX Sequence 451 AA:

alignment_scores: Quality: 169.50 Length: 483
Ratio: 0.712 Gaps: 21
Percent Similarity: 49.275 Percent Identity: 20.290

alignment_block:
US-09-303-518D-125 x AAY34343 ..

Align seg 1/1 to: AAY34343 from: 1 to: 451

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27 AlaGlySProValGluValLeu.....ProLeProSerGlnVal 40
87 CGCGTTCGTCGAGAGATATGCCGGATCGCCCTCGATGAAGTCA 136
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
40 ValIleProLeuGlnGlnHisIleGlyAlaProAlaThrAlaThrVal 57
137 AGGAAGGCGATGCGTCACAAAGAGCCAAAGTGTGTTGAAGACAAAG 186
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57 ySLySgIAspBeluValIlyValGlyThrIleIle.....68
187 AATCCGGGCGTGTGTTTACTGCGCGGCTTACGCAAAATCGCCGGC 234
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69 .....AlaGlnAlaGlyGlyPheValSerAlaAs 78
235 .ATTACCCGTGGC...GAAAAGCCGCTCTTCAG.....264
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78 nIleHisSerSerValSerGlyValLeuLysIleAspAsnValIlyrA 95
265 .....TCAGTGTGATTCGCGTGAAGGCAAC 291
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
95 spSerSerGlyTyrProLysProAlaValPheIleSerValGluGlyasp 111
292 GAGCAATCGAGTTTGAACGCTACGCACTGAGCGCTG.....GCAAA 335
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112 GluTrpGluGluGlyIleAspArgSerProAlaIleValIlySgIucySAs 128
336 CTTAAGCGGCGAAGAGTGGCGCGCAACCTGATCCATCCGATTTGTGA 385
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128 nIleuAspAlaLysGlnIleValAlaLysIleSerAlaIleGlyIle..V 144
386 CTGCGCTGGCGACCCGTCGCTTC.....AGCAAAATT 417
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
144 aIGLyLeuGlyGlyAlaThrPheProThrHisValIlyLeuSerProPro 160
418 CCGCGCTGATGCGGAGCCGCTTCCATCTTCGATTCGATGGAAC 467
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
161 ProGlyAsnLysIleGlu.....IleLeuIleIleAsnAlaValGlyIcy 175
468 CATCGCGTGGCGCTGCGACCTACGATTCATCAAGAGCGCGGAGG 517
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518 ATTTCAAAGCGGCGCTGTGATTGAGCGCTTGAACGCAAAATC 567
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192 IuIleMetIleGlyValSerIleLeuMet.....LysAlaIle 204
568 CATGTTTGAAGGAGCT...GGCGAGAGCGTCCGCTGAAATGCT.. 612
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205 GlnValAsnLysAlaValIleGlyValGluAsnAsnLysLysAspAlaI 221

```

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613 .....GCCAACATCGAACAACATG 631
221 eAlaHisLeuThrIlySleAlaIleThrAlaIlyrProGlyIleGluValMetP 238
632 AATTCGGCGCGCGCATCTCGCGGTTGAGTGGCAGCACCATTCATTC 681
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238 roLeuLysValGlnIlyrProGlnGlyGlyGlnLysGlnLeuIleAspAla 254
682 ATC.....GAGCGGTGGCGCGCAATTA 704
255 ValIleArgLysGlnValIlySerGlyAlaLeuProIleSerThrGlyAl 271
705 AACCGTGTGACCAATCATATATCAAGATGTAATTCATTCGCGCTGTT 754
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271 aValVal.....GlnAsnValGlyThrValPheAlaValT 283
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283 yrcIuAlaValaGlnLysAsnLysProLeuValGluArgIleValThrVal 299
796 GGTGTTCTCAAGTCAACAACCGCGCTTCGCGTACCGTTTGGGTGC 845
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300 ThrGlyLysLysLeuSerArgProSerAsnLeuValArgIleGlyTh 316
846 GAAGTATCGCAATT.....ACTGCGGCGCAATTTGTTGACAGACAGA 889
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316 rProIleAlaIleAlaLeuIleGluAlaIleGlyLeuProGluAsnThrG 333
890 ACCCGTGTATTCGCTTCGCTGATTTGACGCGCGCATTCACAAAGCGCG 939
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333 LysIleIleGlyGlyGly.....339
940 CAGGATTTATTTGGGAGCGTACCAATCAGATTTCCGTTATGGAAGAAG 989
339 .....339
990 CCGCAGCAAGAGCTGTGCGTGGGTGGCGCGCGCACCGCAATTAATCT 1039
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340 .....PrometMetGlyArgAlaLeuLeuSerProAsp...ValP 352
1040 CCATACGCGGTACACACCTGCGCATTTCTCTGAAAACAAACACTCTTCAAG 1089
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352 roValThrIlySgIysSerSerGlyValLeuIleuAspArg.....365
1090 TTCACACAGCGCGTCAACGCGCGGCGCGCATGTCGCGCATTTGTAC 1139
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366 ...GluIleuAlaValaLysIlyPrometAlaArgAspCysIleArgCysAlaIy 381
1140 TTACGAGCGCGGTATGCGCTTGGATATCTGCGCCACCTGCTTTGCGCG 1189
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381 scYsValGlyValCysProMetGlyLeuAsnProAlaPheLeuMetArgA 398
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415 CysIleGluIlySgIysSer.....CysSerPheThrCysPr 426
1281 GGGCAATATACGATACGCGCGCGCTGTCGCAAGTCTGGAACATCT 1329
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426 oAlaAsnArgProLeuLeuAspTyrIleArgGlnAlaLysThrVal 1442

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seq_name: /STDS1/gcgdata/geneseq/geneseqpr_emb1/AA2001.DAT.AAB59813

seq_documentation_block:

ID AAB59813 standard; Protein; 1017 AA.

XX AAB59813;

XX AC AAB59813;

XX DT

XX 04-APR-2001 (first entry)

DE Ttld protein #4.
 XX
 KW Toluene degradation; enzyme; waste degradation; Ttld.
 OS Thauera aromatica.
 OS Xanthomonas maltophilia.
 OS Geobacter metallireducens.
 OS Azarcus toluilyticus.
 XX
 PN W0200072650-A2.
 XX
 PD 07-DEC-2000.
 XX
 PF 24-MAY-2000; 2000MO-US14298.
 XX
 PR 01-JUN-1999; 99US-0323872.
 XX
 PA (UYOH-) UNIV OHIO.
 PI Coschigano PW;
 DR WPI: 2001-041080/05.
 DR N-PSDB; AAF23625, AAF23627.
 XX
 PS Disclosure; Fig 5; 122pp; English.
 XX
 CC The present invention relates to toluene degrading enzyme genes and
 CC proteins tutt (see AAF23629 and AAB59813), tutt (AAF23630 and AAB59832),
 CC tutt (AAF23631 and AAB59833) and tutt (AAF23632 and AAB59834). The
 CC toluene degrading enzymes are homologues of pyruvate formate lyase. The
 CC toluene degrading enzymes are useful for biological treatment of organic
 CC compounds and in particular for the degradation of toluene and its
 CC analogs contained in liquid or solid waste source. The present sequence
 CC is a protein sequence for toluene degrading enzyme, tutt.
 XX
 SO Sequence 1017 AA;

alignment_scores:
 Quality: 166.50 Length: 486
 Ratio: 0.733 Gaps: 26
 Percent Similarity: 46.708 Percent Identity: 25.103

alignment_block:
 US-09-303-518D-125 x AAB59813 ..

Align seg 1/1 to: AAB59813 from: 1 to: 1017

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103 GAATATGCGCGGTATGGCCCTCGATGAAGTAAGAGAGCGATCGCT 152
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125 GATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 141
153 CAAAAAAGGCAAGTGTCTTTGAGACAAAAAAGATCGGGGCGTGT 202
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142 SerProAlaArgSerAlaGlyArgAlaGlyArgAlaGly..... 155
203 TTATCGCGCGCGCTTCAGCAAAATCGCGCATTTCA...CCGTGCGAA 249
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156 ....CysAla.....ArgSerSerArgLysThrSerArgProIleArg 169
250 AACCGCT.....ACTTACGCTAGT..... 269
169 erAlaArgProSerCysSerLysSerProThrSerAlaSerAlaPro 185
270 .....CGTATGCGCGTTGAGACGACGACGAAATCGATTTG 307

```

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186 ProSerProAlaArgAlaSerArgThrArgCysArgArgAsnSerLeuPr 202
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202 oSerSerAlaThrArgSerSerAla.....ThrAlaAlaAlaThrP 216
358 CGCAACCTGATCCTATCGCTTTGAGACTGCGTGGCGACCGCTCGCT 407
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216 roArgArg..LysThrProCysCysLysArgThrThrArgProProSer 232
408 CACCAAAATTCCTCGCGCTGATGCGCGCTTCGCCATCTTGCAATG 457
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232 erThrArg.....AsnSerSerArgAlaThrThrPmet 242
458 CGATGACACCAATCGCTG.....CTGGACACCTACGCTCAT 498
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243 ArgTTrpAsnSerSerArgTTrpAsnValArgPheProSerMetAlaProAl 259
499 ATCAAGAAGACCGCGAGATTTCAAACGCGCTGTGTGATTCAGCCG 548
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259 aserArgAlaProThrAlaLysSerSerArgLysArg.....Thri 273
549 TTGACCGACGCAAAATCCATGTTTGAAGCAGCTGGCGACGCGC 598
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273 leCysSerSerSerProSerAlaAlaProThrProArgAlaArgThrPro 289
599 CGTCAAAATGCTGCAACATCGAACAACATGATTCGGCGCGCGCAT 648
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290 AlaThrThrProThrProSerSerArgLysProSerGlySerAlaAlaPr 306
649 CCGTCCCGCTGTTAGTGGACGACGACATTCATTCATCGAGC..... 688
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689 ..CGTGGCGCGGATTAACCGCTGACCATCATTAATCAAGATGTA 736
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316 rArgThrAlaArgArgArgCysAlaGly..... 325
737 TTACATTGGCGCTGTTGTTGCAACGCGCTGCAACCGCGCGGTG 786
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326 .....PheSerSerAlaSerAlaThrAspSerAla.. 335
787 ATTGCCCTAGGTGTTCTCAAGTCAACAACCGCGCTCTTGCTGCTACGT 836
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336 .....IleArgArgSerSerThrThrArgSerAla..... 345
837 TTTGGTGGCAAGATATCGCAATTAATCTGCGCGCATTTGTTGACACG 886
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346 .....ArgSerArgArgAsnThrProSerSerAlaSerThrAla 358
887 ACAACCGCGTATTTCCGCTCGGATTTGACG.....GC 921
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359 ThrAlaPro.....ProThrArgLysProThrThrLysThrCysCys 373
922 GCGATTACACAAGCGCGCGCATTTATTTGGACGTACCAACATTCAGAT 971
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
373 sAlaCysArgProAlaSerThrVal.....AlaAlaArgArgLysL 387
972 TTCCGTTATCGAAGAGCGCGCACCAAGAGCTTTGGCTGGGTCCGC 1021
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387 yspProVal..ArgLysValAlaAlaGlnSerSerArgProSerCysTrp 402
1022 CGCAGCGCGACA.....AATTCGCA..... 1042
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
403 LysSerArgSerMetThrAlaThrThrGlyArgThrProThrCysAsn 419
1043 .....TCACGCGTACAACTTCGCGCATTTCCGTA 1073
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
419 rAlaArgArgProValIleSerArgArgSerProSerArgMetPheGly 436
1074 AAACAACCTCTTCAAGTTCAACACGCGCTCAACGCGCGGACGCGCA 1123
||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

```

436 rgleuSerAlaSerSerIleasMetarGerThrsValSerAlaPro 452
 1124 ...TGTCGGCATTTGACTTACGACGGCGTATGCCCTTGATATCCTG 1170
 453 ArgThrcysAlaThrSerSerSerAlaSerCysArgCysLeuSerCys 469
 1171 CCCA.....CCCCGCTTTT 1184
 469 sProGlnSerThrThrAlaAlaThrPasnSerGlyTTPThrProAlaProC 486
 1185 GCGGATTTAATCGTCGCGATACCGACGCGCGCAT.....1225
 486 ysProSerSerProMetAlaGlyThrThrArgSerArgArgSerSerArg 502
 1226TGGTTGCTTGGAATTTGACGACGAAGAAGACCTCGCTTG 1263
 503 ArgThrProSerThrProSerArgAsnTrpTyrSerArgArgArgAsnTh 519
 1264 TGCAGCTTCGTCGCGCGCAATACGAATACGCGCGCTGTCGCA 1313
 519 rProSerSerAsnSerAlaLysArgArgGlyThrGlyLysValSerArgL 536
 1314 AGTGC 1318
 536 yscys 537
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 AC AAB59826;
 XX
 DT 04-APR-2001 (first entry)
 XX
 DE Protein #3 encoded by Tuld/E gene.
 XX
 KM Toluene degradation; enzyme; waste degradation; Tule; Tuld.
 XX
 OS Thauera aromatica.
 OS Xanthomonas maltophilia.
 OS Geobacter metallireducens.
 OS Azococcus toluyticus.
 XX
 PN W0200072650-A2.
 XX
 PD 07-DEC-2000.
 XX
 PF 24-MAY-2000; 2000MO-US14298.
 XX
 PR 01-JUN-1999; 99US-0323872.
 XX
 PA (UYOH-) UNITV OHIO.
 XX
 PI Coschigano FW;
 XX
 DR WPI: 2001-041080/05.
 DR N-PSDB; AAF23627.
 XX
 PT Composition comprising toluene degrading enzyme useful for biological
 PT treatment of organic compounds, especially for degrading toluene or its
 PT analogs
 XX
 PS Disclosure; Fig 12; 122pp; English.
 XX
 CC The present invention relates to toluene degrading enzyme genes and
 CC proteins tuch (see AAF23629 and AAB59831), tult (AAF23630 and AAB59832),
 CC tulf (AAF23631 and AAB59833) and tutt (AAF23632 and AAB59834). The
 CC toluene degrading enzymes are homologues of pyruvate formate lyase. The
 CC toluene degrading enzymes are useful for biological treatment of organic
 CC compounds and in particular for the degradation of toluene and its
 CC analogs contained in liquid or solid waste source. The present sequence
 CC is a protein sequence encoded by toluene degrading enzyme gene, Tuld/E.

XX
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 alignment_scores:
 Quality: 166.50 Length: 486
 Ratio: 0.733 Gaps: 26
 Percent Similarity: 46.708 Percent Identity: 25.103
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 710 LysProArgProThrCysArgProSerProGlyThr.....AlaAr 723
 103 GAATATGCGGTATGCGCCCTCGATGAAAGTCAGGAAGGCGATCGGT 152
 723 garGValSerThrThrSerProArg..SerThrGlyArgArgTrpSer 739
 153 CAAAAAAGGCCAAGTCTGTTGAAGACAAAAAGATCGGCGGTGT 202
 740 SerProAlaArgArgSerAlaGlyArgAlaGlyArgAlaGly..... 753
 203 TTATCGCGCGGCTTCAGGCAAAATGCGCGATTCGA...CGTGGCGAA 249
 754 ...CysAla.....ArgSerSerArgLysThrSerArgProIleArgS 767
 250 AAGCGCGT.....ACTTCAGTCAGT..... 269
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 784 ProSerProAlaArgAlaSerArgThrArgCysArgArgAsnSerLeuPr 800
 308 AACGTAACGACCTGAAGCGCGTGGCAAACTTAAGCGCGCAAGAAGTGC 357
 800 oSerSerValThrArgSerSerAla.....ThrArgAlaAlaThrP 814
 358 CGCAACCTGATCCAAATCGGTTGTGACTGCGTCGCGACCCGTCGTT 407
 814 roArgArg..LysThrProCysCysLysArgThrThrArgProprosers 830
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 830 erThrArg.....AsnSerSerArgAlaThrTrpMet 840
 458 CGATGACACCAATCCGCTGG.....CTGCGACCCCTACGGTCATT 498
 841 ArgThrPasnSerSerArgTrpAsnValArgPheProSerMetAlaProl 857
 499 ATCAAGAGACGGCGCGAGATTTCAAACGGCGCTTGTGATATGACCG 548
 857 aserArgAlaProThrAlaLysSerSerArgLysArg.....ThrI 871
 549 TTTGACCGACGCAAAATCCATGTTGTAAAGCAGCTGGCGCGACAGTCG 598
 871 leCysSerSerSerProSerAlaAlaProThrProArgAlaArgThrPro 887
 599 CGTCTGAATAATGCTGCACATCGAAGCAACATGAATTCGGCGCGCCGAT 648
 888 AlaThrThrProThrProSerSerArgGlnProSerGlySerLalaArgPr 904
 649 CTGCGCGGTTGATGTCGACGACCATTCATTATGTCGAC..... 688
 904 oSerProPro.....SerSerSerAlaIleProA 914
 689 ..CGTGGCGCGCAATTAACCGTGGACCATTCATTAACAGATGTAA 736
 914 rGArgThrAlaArgArgArgCysAlaGly..... 923

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737 TTACATTGGCCGTTTGTTCACAGCCGCTTGACACCGAGCGCTG 786
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XX      : : : : : : : : : : : : : : : : : : : : : :
XX      : : : : : : : : : : : : : : : : : : : : : :
934      : : : : : : : : : : : : : : : : : : : : : :
XX      : : : : : : : : : : : : : : : : : : : : : :
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XX      : : : : : : : : : : : : : : : : : : : : : :
944      : : : : : : : : : : : : : : : : : : : : : :
XX      : : : : : : : : : : : : : : : : : : : : : :
887 ACACCGCGGTGATTTCCGTTTCGATTGAAAGC.....GC 921
XX      : : : : : : : : : : : : : : : : : : : : : :
957 ThrAlaPro.....ProThrArgLysProThrThrGlySerThrCys 971
XX      : : : : : : : : : : : : : : : : : : : : : :
922 GCGATTACACAAGCGCGCGCATTTATTTGGACGCTACCAATCAGAT 971
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971 sAlaCysArgProAlaSerThrVal.....AlaAlaArgArgLysL 985
XX      : : : : : : : : : : : : : : : : : : : : : :
972 TTCGCTATTCAGAGAGCGCGCAAGACGCTGTTCGGCTGGGTGGC 1021
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XX      : : : : : : : : : : : : : : : : : : : : : :
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1084 ySProSerSerProMetAlaGlyThrThrArgSerArgArgSerArg 1100
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1226 .....TGGTTCCTTGGATTTGACGAGACAGACCTCGCTTTG 1263
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XX      : : : : : : : : : : : : : : : : : : : : : :
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XX      : : : : : : : : : : : : : : : : : : : : : :
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ID   AA104998 standard; Protein; 388 AA.
AC   AA104998;
XX
XX   06-JUL-1999 (first entry)
XX
XX   Mycobacterium species protein sequence 50B.
XX
XX   Secreted protein; Mycobacterium; primer; PCR; amplification; probe.
XX

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```

KW      hybridisation; detection; vaccine; immunisation; infection.
XX
XX      Mycobacterium sp.
XX
XX      W0909186-A2.
XX
XX      25-FEB-1999.
XX
XX      14-AUG-1998; 98WO-FR01813.
XX
XX      11-SEP-1997; 97FR-0011325.
XX      14-AUG-1997; 97FR-0010404.
XX
XX      (INSP ) INST PASTEUR.
XX
XX      Gicquel B, Lim EM, Pellicle V, Portnoi D, Gouget de la Salmoniere Y,
XX      Guigueno A;
XX      WPI; 1999-181045/15.
XX      N-PSDB; AAX34249.
XX
XX      Mycobacterial DNA vectors containing reporter constructs - for
XX      identifying coding or promoter sequences involved in
XX      infection-associated protein expression
XX
XX      Claim 32; Fig 50B; 309pp; French.
XX
XX      Sequences AA104742-Y05000 and AA107201-Y07204 represent secreted
XX      proteins from various Mycobacterium species microorganisms. The
XX      encoding nucleotide sequences can be used as primers and probes for
XX      methods for detecting and identifying mycobacteria, especially belonging
XX      to the M. tuberculosis complex. The encoded proteins can be used in
XX      vaccines for immunisation against a bacterial or viral infection.
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XX      Quality: 148.00      Length: 490
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18 AGGTCTAAACCTGCCCCATCGCGGCGACGCGAGCAGCAGCCGTTTACGACG 67
XX      : : : : : : : : : : : : : : : : : : : : : :
13 ArgSerArgArgCysGlyArgCysArgArgAlaValGlyArgArgArgAr 29
XX      : : : : : : : : : : : : : : : : : : : : : :
68 GCCCGCGCATTAACGCAAGTCG.....G 90
XX      : : : : : : : : : : : : : : : : : : : : : :
29 gArgTrpGlnTrpArgGlnArgArgArgCysGlnHisArgGlnArgTrpA 46
XX      : : : : : : : : : : : : : : : : : : : : : :
91 TTGCTTGGCGAAGATATCGCGGTAT.....GCGCCCTCGATGAA 131
XX      : : : : : : : : : : : : : : : : : : : : : :
46 rArgTrpProArgArgCysArgTrpArgTrpArgMetAlaLeuArgArg 62
XX      : : : : : : : : : : : : : : : : : : : : : :
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63 ArgArgGlyArgArgTrpArgArgThrArgArgAsnArgProArgArgAr 79
XX      : : : : : : : : : : : : : : : : : : : : : :
182 AAAAGATCCGCGCGGTGGTTTACTGCGCGCGCTTCAGCGCAATCGCC 231
XX      : : : : : : : : : : : : : : : : : : : : : :
79 gArgArgArgArgArgArgGly.....ProGlyArgArgArgProArgTr 93
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232 GCGATTCAACCGTGGCGAAGACGCGTACT..... 260
XX      : : : : : : : : : : : : : : : : : : : : : :
93 hrValGlyTrpArgArgArgArgTrpArgArgAlaArgArgTrpTyr 109
XX      : : : : : : : : : : : : : : : : : : : : : :
261 .....TCAGTCAGTCGTGATGCCGT..... 281
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282 .....TGAAAGCAGACGACCAATCGAGTTTGAAACGCTACGACACT 321
126 GArgTrpAlaAspArgGlnArgArgGly..... 135
322 GAAGGGCTGGCAAACTTAAGCGGGGAGAAAGTAGCGCCGCAACCTGATCA 371
136 .....ArgArgArg..... 138
372 ATCCGGTTTGTGACTGCGCTGCCACCCGCTCCGTTGACGAAATTCCTG 421
138 ..... 138
422 CCGTCGATGCCGAGCCGTCGCAATCTTCGTAATGCGATGACACCAAT 471
139 .ArgArgCysArgArgValArgArg.....GlyArgArgS 150
472 CCGCTGGCTGGCCAGCCCTGATCAAGAGAGCCGCGAGATTT 521
150 eArgArgCysArgArgGlnArgHisAlaAspArgAlaArgArgArg 166
522 CAACGCGCCCTGTGTATGAGCCGTTGACGCAACGCAAAATCCATG 571
167 ArgArgArg.....ArgArgGlnGln..... 173
572 TTTGTAAGCAGCTGGCCAGACGTCGCTGAAATGC.....TGC 615
174 ...PheArg**TrpArgGlyArgArgArgGlyArgCysArgArgAla 189
616 AACATCGAAGACATGAAATTCGCGCGCCGATCCTGCGGTTTGAGTGG 665
189 rGlnArgGlnTrpArgValArgArgProArgArgSerArgHis..... 203
666 CAGCAGCATTCATTCATCGAGCCGCTGCGCGCAATAAACCCTGTGGA 715
204 .....GlyArgGlnHis..... 207
716 CCATGATTAATCAATGTAATTAATTAATGCGGTTGTTGCAACAGGC 765
208 .....ArgTrpArgTrpArg.....ArgArg 215
766 CGTGTGAACCCGAGCGGCTGATGCGCTAGGTGTTCTCAATCAACAA 815
215 rArgTrpArgGlnAlaAspArgProArgTrp..... 226
816 ACCGGCCCTTTCGCTACCGCTTTGGGTGCGAAAGTATCGCAATCTACTG 865
227 .....ArgArgArgCys..... 230
866 CCGGCGAATTTGGTGTACAGACAAACCGGCTGATTCGCGTTCCGTATTG 915
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966 TCAGATTCCTGATTCGAAGAAAGCCGACGCAAGA..... 1001
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1002 ....GCTGTGGCTGGTGGTGTG.....GCCGAGAGCC 1028
272 yGlyGlyArgGlyGlyGlyAspGlyPheValArgCysTrpArgProTh 289
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289 rGlnArgHisArgProThrArgLeuAlaPro.....ile 299
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300 AsnGlnGlyPheGlyAlaGlyProGlnHisGlyHisProLeuSerTrpArg 316

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316 gSerGlyLeuArgTrpCysArgGlySileu..... 326
1161 GATATCTCTGCCACCCCTCTTGGCGATTTAAATCGTGGCATACCG 1210
327 .....ProGlnArgLeuArgGlyGlnPheLeuAla 339
1211 ACAGCCGCGAGGATTTGGTCTTGAATTTGACAGCAAGAACCTCGCT 1260
340 Thr.....ProThrGlnG 344
1261 TTGTGCGATTCGTCGCCGCGCAATACGATACG...GCCGCTGT 1307
344 vAlaTrnGly**AlaIleArgAlaAlaHisSerGlyGlyHisGlyC 361
1308 GCGCAAAAGTCTGGAACCA 1327
361 ys**AlaCysTrpLeuPro 367

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ID AAB59827 standard; Protein; 1592 AA.
AC AAB59827;
XX
XX
XX 04-APR-2001 (first entry)
XX
XX Protein #4 encoded by TufD/E gene.
XX
XX Toluene degradation; enzyme; waste degradation; TufE; TufD.
XX
XX Thauera aromatica.
XX
XX Xanthomonas maltophilia.
XX
XX Geobacter metallireducens.
XX
XX Azotobacter vinelandii.
XX
XX WO200072650-A2.
XX
XX 07-DEC-2000.
XX
XX 24-MAY-2000; 2000WO-US14298.
XX
XX 01-JUN-1999; 99US-0323872.
XX
XX (UYOH-) UNIV OHIO.
XX
XX Coschigano PW;
XX
XX WPT: 2001-041080/05.
XX
XX N-PSDB; AAF23627.
XX
XX Composition comprising toluene degrading enzyme useful for biological
XX treatment of organic compounds, especially for degrading toluene or its
XX analogs
XX
XX Disclosure; Fig 12; 122pp; English.
XX
XX The present invention relates to toluene degrading enzyme genes and
XX CC proteins tufH (see AAF23629 and AAB59831), tufI (AAF23630 and AAB59832),
XX CC tufF (AAF23631 and AAB59833) and tufG (AAF23632 and AAB59834). The
XX CC toluene degrading enzymes are homologues of pyruvate formate lyase. The
XX CC toluene degrading enzymes are useful for biological treatment of organic
XX CC compounds and in particular for the degradation of toluene and its
XX CC analogs contained in liquid or solid waste source. The present sequence
XX CC is a protein sequence encoded by toluene degrading enzyme gene, TufD/E.
XX
XX Sequence 1592 AA:

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Quality: 146.00 Length: 527
Ratio: 0.705 Gaps: 31
Percent Similarity: 39.279 Percent Identity: 23.909

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US-09-303-518d-125 x AAB59827 ..

Align seg 1/1 to: AAB59827 from: 1 to: 1592

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689 ArgAlaIaIaProIaIaCysAlaIaArgPheAlaIaArgLeuGlyPro 705
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63 .CGAGCGCCCGGCGCATTCACGAGTCGGCTTGGCGGCAAGAAATATGCC 111
   ||||| ||||| ||||| ||||| |||||
705 OArgGlyThrSerArgGlyArgSerArgCysSerProAspArgCysCys 722
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112 GGTATGCGCCCGCTCGATGAAGTCAAGAGCGCATGCCGTCAAAAAGG 161
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722 IArgTrpSerArgCysSerProSerProArgArgCysProProSerSer 738
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152 CCAAGTGGCTGTTGAAGCAAAAATCCGCGCGCTGTTACTGCGC 211
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739 ProAla.....GlyAlaProGlyAlaIaIaCys.. 747
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212 CGGCTTCAGGCAAAATCCGCGCATTCACCGTGGCAAAACGGCTACTT 261
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748 .....SerArgArg.....ProPheSerArgSerArgAspS 758
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262 CAGTAACT..CGTATGCGCGCTGAAGGCAACGCAATTCGATTTGA 308
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758 erAlaGlyProArgAlaIaIaArgPheArgArgCysArgAsp..... 771
   ||||| ||||| ||||| ||||| |||||
309 ACGTACGACACTGAAGCGCTGGCAAACTTAAGCGGCAAGAAAGTGGCC 358
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772 .....AlaCysGlyIaIaArgAlaIaIaArgCysPro 780
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359 GCAACTGATCCAAATCCGCTTGTGAGTCGCGTCCGACCCGTCCTC 408
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780 OGlyPro.....ArgSerAlaProSerIleA 789
   ||||| ||||| ||||| ||||| |||||
409 AGCAAAATTCCTGCCGTCGATGCCGAGCCGTCGC..... 443
   ||||| ||||| ||||| ||||| |||||
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444 .....CATCTGTCATATGCATG 463
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806 LeuCysGlyAlaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 482
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464 ACACCAATCCGCTGCTGCGACCGCTACGGTCATTCAAAGAACG.... 509
   ||||| ||||| ||||| ||||| |||||
822 eGlyAlaSerSerGlyCys...ProHisProValaIaIaIaIaIaIaIa 838
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510 .....CGCCGAGGATTT 521
   ||||| ||||| ||||| ||||| |||||
838 alaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 853
   ||||| ||||| ||||| ||||| |||||
522 CAAACGCGCGCTGTTGATTAAGCCGTTGACCGAGCAAGCAAAATCAG 571
   ||||| ||||| ||||| ||||| |||||
854 .....ArgPheArgGlyProIaIaIaIaIaIaIaIaIaIaIaIaIa 868
   ||||| ||||| ||||| ||||| |||||
572 TTTTGAAGGAGCTGGGCGACAGCTGCCGTCTGAATAATGC...TGCCA 618
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868 s.....TrpArgTrpProArgProArgArgCysArgCysSerA 881
   ||||| ||||| ||||| ||||| |||||
619 ATCGAAACACATGAATTCGGCGGCC.....GCATCTGC...CGG 656
   ||||| ||||| ||||| ||||| |||||
881 rArg.....TrpGlyArgProIaIaIaIaIaIaIaIaIaIaIaIaIa 894
   ||||| ||||| ||||| ||||| |||||
657 TTTGAGTGGAC.....GC 670
   ||||| ||||| ||||| ||||| |||||
895 AlaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 911
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721 AATTATCAAGATGTAATTACCATTTGGCCGTTGTTGGCAACAGCCGCT 770
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928 rArgArgSerArgCys.....ProIaIaSerProIle 938
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939 ArgTrpIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 949
   ||||| ||||| ||||| ||||| |||||
821 GCCTTTGCGCTACCGTTTGGCTGGCAAGTATCGCAAT.....T 861
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949 gProIaIa.....GlyCysSerProArgAlaIaIaIaIaIaIaIaIa 962
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862 ACTCGCGCGCAATTTGTT.....TGACACAGA 887
   ||||| ||||| ||||| ||||| |||||
962 rGcysGlyIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 978
   ||||| ||||| ||||| ||||| |||||
888 CAACCGCGT.....GATTTCCGGTTCGTTATTTGAACGCGC 922
   ||||| ||||| ||||| ||||| |||||
979 SerIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 994
   ||||| ||||| ||||| ||||| |||||
923 CGATTACAAAGCGCGCGCAAGTATTTGGAGCGCTACCAATCAGATT 972
   ||||| ||||| ||||| ||||| |||||
994 gSerThrIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 1011
   ||||| ||||| ||||| ||||| |||||
973 TCCGTTATCGAAGCGCGCGCAAGAGCTTTGCGCTGGGCTGGCC 1022
   ||||| ||||| ||||| ||||| |||||
1011 hIaIaSerThrIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 1025
   ||||| ||||| ||||| ||||| |||||
1023 .....GCAGCCGAC 1033
   ||||| ||||| ||||| ||||| |||||
1026 ThrAspIleHisSerGlyIaIaIaIaIaIaIaIaIaIaIaIaIaIa 1042
   ||||| ||||| ||||| ||||| |||||
1034 AATATCTCATCAAGCGCTACACCTCGGCCATTTCTGTAACAAACATC 1083
   ||||| ||||| ||||| ||||| |||||
1042 gAlaIaIaSerGlyAlaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 1059
   ||||| ||||| ||||| ||||| |||||
1084 TTCAGTTCACACACAGCGCTCAACG..... 1109
   ||||| ||||| ||||| ||||| |||||
1059 ysProValaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 1075
   ||||| ||||| ||||| ||||| |||||
1110 .....CGGCGACCGCGCCATGTCGCGCATGTTGTTACTT 1141
   ||||| ||||| ||||| ||||| |||||
1076 AlaCysGlySerSerSerArgArgProSerSerGlyIaIaIaIaIaIa 1092
   ||||| ||||| ||||| ||||| |||||
1142 ACGAGCGCTGAT.....GCCCTGGATAT..... 1166
   ||||| ||||| ||||| ||||| |||||
1092 lProIleArgProSerSerIleCysGlyIaIaIaIaIaIaIaIaIaIa 1109
   ||||| ||||| ||||| ||||| |||||
1167 .....CTGCCCCAC 1175
   ||||| ||||| ||||| ||||| |||||
1109 roSerSerProIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIaIa 1125
   ||||| ||||| ||||| ||||| |||||
1176 CCGTCTTTTGGCGATTAATGTCGCGCATACCAACGCGCGAGGAT 1225
   ||||| ||||| ||||| ||||| |||||
1126 ThrProCysArgArgHisAsnArgArgArgGly..... 1136
   ||||| ||||| ||||| ||||| |||||
1226 TGGGTTCTTGGAAATTTGACGAGAAGACCT 1256
   ||||| ||||| ||||| ||||| |||||
1137 .....GlySerArgArgPro 1141
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seq_name: /SISL/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:AAB59817

seq_documentation_block:

ID AAB59817 standard; Protein: 999 AA.

XX AAB59817;

XX AC

DT 04-APR-2001 (first entry)

XX

DE Tutd protein #8.
 KM Toluene degradation; enzyme; waste degradation; Tutd.
 XX
 OS Thauera aromatica.
 OS Xanthomonas maltophilia.
 OS Geobacter metallireducens.
 OS Azotobacter toluyticus.
 XX
 PN W0200072650-A2.
 PD
 PD 07-DEC-2000.
 PF 24-MAY-2000; 2000MO-US14298.
 PR 01-JUN-1999; 9905-0323872.
 PA (UYOH-) UNIV OHIO.
 PI Coschignano PW;
 DR WPI; 2001-041080/05.
 DR N-PSDB; AAF23625, AAF23627.
 XX
 PT Composition comprising toluene degrading enzyme useful for biological treatment of organic compounds, especially for degrading toluene or its analogs
 PS Disclosure; Fig 5; 122pp; English.
 XX
 CC The present invention relates to toluene degrading enzyme genes and proteins tutd (see AAF23629 and AAB59831), tuti (AAF23630 and AAB59832), CC tutf (AAF23631 and AAB59833) and tutg (AAF23632 and AAB59834). The CC toluene degrading enzymes are homologues of pyruvate formate lyase. The CC toluene degrading enzymes are useful for biological treatment of organic compounds and in particular for the degradation of toluene and its CC analogs contained in liquid or solid waste source. The present sequence is a protein sequence for toluene degrading enzyme, Tutd.
 CC
 SO Sequence 999 AA;

alignment_scores:
 Quality: 145.00 Length: 529
 Ratio: 0.700 Gaps: 31
 Percent Similarity: 39.130 Percent Identity: 23.819

alignment_block:
 US-09-303-518D-125 x AAB59817 ..

Align seg 1/1 to: AAB59817 from: 1 to: 999

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18 AGGTCTAAACCTGCCATCGCGGACAGCCGAGCAAGCCGTTTAA..... 62
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
94 ATGAlaAlaProAlaCysAlaArgArgPheAlaTyrArgArgLeuGlyPr 110
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
63 CGAGCGCGCGCGCATACGGAAGTCGGTTCGCGCGGAGAGAAATATGCC 111
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
110 atgCysThrSerArgGlyArgSerArgCysSerProAspArgCysCysA 127
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
112 GGTATCGCGCCCTCGATGAAGTCAAGAGCGATCGCGTCAAAAAAGG 161
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
127 rgtPrSerArgCysSerSerProAspArgArgCysProProSerSer 143
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
162 CCAAGTGTCTTTGAAGACAAAAAGATCCGGCGGTGCTTTACTGGCC 211
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
144 ProAla.....GlyAlaProGlyAlaThrCys.. 152
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
212 CGGCTTCAGGCAAAATCGCGGATTCACCGTGGGAGAAAGCGGACTT 261
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
153 .....SerArgArg.....ProPheSerArgSerArgAspS 163
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
262 CAGTCACT..CGTGAATGCGCTTGAAGCAACGACGAAATCGAGTTTGA 308

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|||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
163 erAlaGlyProArgAlaAlaArgPheArgArgCysArgAsp..... 176
309 ACCCTACGCAACCTGAAGCGCTGGCAAACTTAAGCGCGGAGAGAGCGGC 358
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
177 .....AlaCysGluAlaArgAlaArgCysPr 185
359 GCACCTGATCCAAATCGGTTGTGTGACCTGCGCTGCGCACCGCTCGTTC 408
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
185 OGlyPro.....ArgSerAlaProSerIleA 194
409 AGCAAAATTCCTGCGCTCGATGCGGAGCCGTTGCG..... 443
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
194 rglArgGlySerArgAspArgSerArgAlaSerArgSerArgSerArgGly 210
444 .....CATCTGCTCAATG 457
211 SerProLeuCysGlyAlaThrAlaThrSerCysProArgArgArgArgCys 227
458 CGATGACACCAATCCGCTGCGCTGCGGACCGCTACGGTCAATTCAAGAA 507
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
227 sSerIleGlyAlaSerSerIleCys..ProHisProProValArgArgS 243
508 GC.....CGCCGA 515
243 erProValaSerSerIlySerArgAlaHisArgCysThrAlaArgArg 259
516 GGAATTCAAACGGCGGCGCTGTGGTATGACCGCTTACCGAGACGCAAA 565
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
260 Gly.....ArgPheArgGlyProThrSerArgAspThrIlyArgArg 273
566 TCCATGTTTGTAAAGCAGCTGCGGCGCAGACGTCGCGTCAAAATGC...T 612
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
273 gArgCys.....TPrArgTrProArgProArgArgCysArgC 286
613 GCCAACATTCGAACACATGATTCGCGCGCC.....GCATCTCG 653
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
286 ySerArgArg.....TPrIlyArgProLeuThrAlaSerGlyCys 299
654 ...CGTTTGAGTGCAC..... 668
300 ProArgAlaArgTrPrArgArgGlySerAsnTrpSerSerIlyArgSer 316
669 ....GCACATTCATTCATCGAGCGCGCTGCGCGGAGTAATAACCGTGTG 714
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
316 rAlaAlaSerProIlySerArgThrCysGlyArgValArgSerAspHis 333
715 ACATCAATTAATCAAGATTAATTACCATTCGCGCTGTTTGCACACAG 764
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
333 erAlaArgArgSerArgCys.....ProAlaSerSer 343
765 CCGTCTGAACACGAGCGCGTATGCGCTAGTTCGTTCAAGTCAACA 814
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
344 ProIleArgTrPrThrGlyArg...CysArgArgTrp..... 354
815 AACCGCGCTCTGCTGACGAGCTTTGGTGGCGAAGATATGCAAT... 860
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
355 ArgArgProLeu.....GlyCysSerProArgAlaThrCysT 367
861 .....TACTGCGGCGAATGT.....TGA 881
367 hrAlaArgCysGlyArgAspIlyCysSerAlaPhePheGlyAsnProLeu 383
882 CACAGACAAACCGCT.....GATTCGCGTTCGGTATTGA 916
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
384 HisArgSerLeuArgGlyProTrPrAlaAlaProPheArg...AlaHisAr 399
917 ACGGCGGATTAACAAGCGCGACGATTAATTTGGACCGTACCAACAAT 966
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
399 gSerArgSerThrThrArgArgCysAlaValArgIlySerSerArgHisA 416
967 CAGATTTCGTTATCGAAGAGCGCGACGCAAAAGCTTCGGCTGGGT 1016
   |||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:

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416 spArgThrAlaSerThrArgArgProHisLysProPro.....LysGly 430
1017 TCGGCC.....GCGGC 1027
431 CysAlaThrAspIleHisSerGlyArgGlyCysTrpProAlaThrAlaSe 447
1028 CGCAAAATACTCCATCACCGGTACACCGGTGGCCATTCTCTGAAAG 1077.
447 rSerArgAlaAlaSerGlyAlaSerAlaLysArgThrArgLeuArgArg 464
1078 AATCCTTCAGTTCACACGCGGTCAACG..... 1109
464 rGserCysProValArgSerProAlaArgArgGlyThrArgAlaAlaTrp 480
1110 .....CGCGACCGCGCGCCATGGTGGCGATG 1135
481 HisSerAlaCysGlySerSerSerArgArgProSerSerGlyAlaArgProTr 497
1136 GTACTTACGACGCGGTGAT.....GCCCTTGATAT. 1166
497 pSerValProIleArgProSerSerIleCysGlyArgAlaValGlyLeuT 514
1167 .....CCT 1169
514 hrSerProSerSerProLeuAsnArgProPheAlaArgArgSerAlaPro 530
1170 GCCCACCTGCTTTGCGCATTTAATCGTCGCGATACCGACAGCGCGC 1219
531 AlaSerThrProCysArgArgHisAsnArgArgArgTyr..... 543
1220 AGGCATTGGTGTCTGGATTTGACGACGAAGACCT 1256
544 .....GlySerArgArgPro 548

seq_name: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:ABG03569
seq_documentation block:
ID ABG03569 standard; Protein; 819 AA.
XX
AC ABG03569;
XX
DT 13-FEB-2002 (first entry)
XX
DE Novel human diagnostic protein #3560.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
XX
DR WPI: 2001-639362/73.
DR N-PSDB: AAS67756.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity
XX
PS Claim 20; SEQ ID NO 33928; 103pp; English.
XX

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CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. ABG00010-ABG30377 represent novel human
 CC diagnostic amino acid sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 819 AA;

alignment_scores:

Quality:	Length:
Ratio: 0.860	Gaps: 26
Percent Similarity: 37.629	Percent Identity: 25.515

alignment_block:

US-09-303-518D-125 x ABG03569 ..

Align seg 1/1 to: ABG03569 from: 1 to: 819

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353 TGGCGCCGACCTGATCCATCCGGTTGTGTGAGCTGCGCTGCGACCCGT 402
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328 CysProAlaLysArgGlyGlnProGlyCysGly***AlaProTr 342
403 CCGTTCAGCAAAATTCCTGCGCGTGCATGCCGCGTCCGATCTTGT 452
    ||| ||| ||| ||| ||| ||| |||
342 pArgPro.....LeuProArgArgProSerSerValProProPro 356
453 CAATGGATGACACCAATCCGCTGCTGCGACCCCTACGGTCATTATCA 502
    :| :| :| :| :| :| :| :| :| :|
356 la.....TrpSerProGlnAsnLeuProLeuGlySerLeuPro 370
503 AAGAACCGCGCGCATTTCAACCGCGCTGTGTGATGACCGGTT. 551
    ||||| :| :| :| :| :| :| :|
371 AlaLysProThrAsnGlyGlyProAlaLeuCysPhe.ProProProHis 387
552 .....GACGACGCAAAAT..... 566
387 erLeuGlnProGlnAspAlaSerGlyLysThrGlnGlyProGluGluAla 403
567 .....CCATGTTTGTAAAGCAGCTG..... 587
404 ProProProCysLeuValProArgTrProProAsnSerAsnSerArg** 420
588 .....CGCAGACGTGCGCTGTGAAATGCT..... 612
420 *HisProArgArgSer.ProMetSerProAlaProHisSerThrProGly 436
613 .....GCCAACATGCAACACATGAAATC..... 636
437 ArgArgHisLeuThrGlnIleProAsnTrpLysThrHisLeuPhePro** 453
637 .....GGCGGCCGATCTGCTG..... 654
453 *AlaProAlaArgGlyProSerProGlyArgAlaCysThrSerProCysp 470
655 .....GGTTTG.....AGTGGC 666
    |||||

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470 roatrginglyleutrrprarprproalaalargalaThrSergly 486
667 .....ACGCACATTCATTTCATCGAGCGCGCGCAATTAACCGCT 710
487 AlaLeuSerHisLeuHisPheProProThrProAla..Leu...ProA 502
711 GTGGACCATCAATTTATCAAGATGTAATTTACCATTTGGCCGTT..... 751
502 lathPheSerLeuSerLeuGlnLeuProLeuHisLeuProHis 518
752 TGTTCGAACAGCGCGCTCTGAACACGAGCGCGGTATGCGCCGTAGTGT 801
519 CysValGlnArgAla.....ProAlaAlaAl 527
802 TCTCAAGTCACAAACCGCGCTCTTGGCTACCGTTTGGGTGCGAAAGT 851
527 alaIaGlySerArgArgSerArgCysProPro..... 538
852 ATCGCAAAATTTACTGCGGCGAATTGTTGACACAGACAACCGCGTATTT 901
538 ..... 538
902 CCGGTTCCGTTATGAAGCGCGGATTACACAAGCGCGCAGCATTTATTTG 951
538 ..... 538
952 GGACGCTACCAACATTCAGATTCCTTATTCAGAAAGCGCGCAAGA 1001
539 .....SerArgArgSerPro.....AlaCysLeuThrS 548
1002 GCTGTTCGGCTGGTGGCGCGCGAGCAAAATCTCATTCACGCGTA 1051
548 exProThrAlaPheMetArgSerProThrSer***ProSerArgGln 564
1052 CAACCC.....TCGGCCATTCCTTAAAAAACAAC..... 1081
565 ProProTrpSerSerAlaSerThrSerLysArgThrSerValSerSe 581
1082 .....TCTCAAGTTCAACACGCGCTCAACGCGCGAGCGCGCAT 1124
581 rtrPalasSerSerProSerProSerProThrLysSergLysrPhePro 598
1125 GGTGC.....CGATTGTACTTACGAGCGCGTATGCCCT..... 1159
598 rplla***Argarg***LysAlaProAlaSerThrCysProArgArgPro 614
1160 .....TCGATATTC.....TG 1170
615 ThrGlyAlaAlaCysCysValAsnTrpArgSerProLysGlyProGlyAr 631
1171 CCCACCCCTGCTTTTGGCGATTAAATCGTCGGGATACCGACA...GCCG 1217
631 gProPro.....GlySerAlaProProThrAlaAlaIag 642
1218 GCAGGCAT 1225
642 lnaRghis 644
seq_name: /SIDS1/gcgdata/geneseq/geneseqp-embL/AA2001.DAT:AA94220
seq_documentation_block:
ID AAB94220 standard; Protein: 639 AA.
AC AAB94220;
XX
XX 26-JUN-2001 (first entry)
XX
XX Human protein sequence SEQ ID NO:14581.
XX
XX Human; primer: detection; diagnosis; antisense therapy; gene therapy.
XX
XX Homo sapiens.
XX

```

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PN EP1074617-A2.
XX
XX PD 07-FEB-2001.
XX
XX PF 28-JUL-2000; 2000EP-0116126.
XX
XX PR 29-JUL-1999; 99JP-0248036.
XX PR 27-AUG-1999; 99JP-0300253.
XX PR 11-JAN-2000; 2000JP-0118776.
XX PR 02-MAY-2000; 2000JP-0185767.
XX PR 09-JUN-2000; 2000JP-0241899.
XX
XX PA (HELI-) HELIX RES INST.
XX
XX PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
XX PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX
XX DR WPI: 2001-318749/34.
XX
XX PS Claim 8; SEQ ID 14581; 2537bp + CD ROW; English.
XX
XX CC The present invention describes primer sets for synthesizing 5602
XX CC full-length cDNAs defined in the specification. Where a primer set
XX CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary
XX CC to the complementary strand of a polynucleotide which comprises one of
XX CC the 5602 nucleotide sequences defined in the specification, where the
XX CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination
XX CC of an oligonucleotide comprising a sequence complementary to the
XX CC complementary strand of a polynucleotide which comprises a 5'-end
XX CC sequence and an oligonucleotide comprising a sequence complementary to a
XX CC polynucleotide which comprises a 3'-end sequence, where the
XX CC oligonucleotide comprises at least 15 nucleotides and the combination of
XX CC the 5'-end sequence/3'-end sequence is selected from those defined in
XX CC the specification. The primer sets can be used in antisense therapy and
XX CC in gene therapy. The primers are useful for synthesizing polynucleotides,
XX CC particularly full-length cDNAs. The primers are also useful for the
XX CC detection and/or diagnosis of the abnormality of the proteins encoded by
XX CC the full-length cDNAs. The primers allow obtaining of the full-length
XX CC cDNAs easily without any specialised methods. AAH01166 to AAH13628 and
XX CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to
XX CC AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632
XX CC represent oligonucleotides, all of which are used in the exemplification
XX CC of the present invention.
XX
XX SQ Sequence 639 AA;
XX
XX
XX alignment_scores:
XX Quality: 119.50 Length: 486
XX Ratio: 0.504 Gaps: 22
XX Percent Similarity: 48.765 Percent Identity: 22.634
XX
XX alignment_block:
XX US-09-303-518D-125/rev x AAB94220 ..
XX
XX Align seg 1/1 to: AAB94220 from: 1 to: 639
XX
XX 1304 AGCGGCGGTATTCGTAATTCGCCGCGGCAACGAGTGTGCACAAAGCAG 1255
XX |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
XX 95 SerAlaProPheThrSerLeuProPheSerThrSer..... 106
XX
XX 1254 GTCCTCTGTCGCCAATTCACAGCAACCAATGCGCGCGCTGTG.... 1209
XX |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
XX 107 .SerSerAlaAlaSerThrSerAsnProAsnSerAlaSerLeuSerSery 123
XX
XX 1208 .....GTATCGCGCAGCATTAATTCGCGCAAA 1182
XX |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
XX 123 alphaIaGlyLeuProLeuProLeuProProThrSergLysLeuSer 139
XX

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1181 AGCAGGGTGGGAGATATCAAGGGCATCAGCGCTGTAGTA...CC 1135
140 AsnProThrProValIleAlaGlySerThrProSerValAlaGlyPyr 156
1134 AATGGCACCATTGGCGGGTGGCCCGCTTGACGGCT..... 1098
156 OleuGly.....ValAsnSerProLeuSerAlaLeuGlyPheL 171
1098 ..... 1098
171 euThrSerAsnSphRAsnLeuIleAsnSerSerAlaLeuSerSerAla 187
1097 .....GTGTTGAACCTTGAAGATTGTTTTCAGGAATGGCC 1060
188 ValThrSerGlyLeuAlaSerLeuSerSerLeuThrLeuGlnAsn.... 202
1059 GAGGGTGTACCGCTGATGGATATTGTCCGGCTGGCGCCACCCAGC 1010
203 .....SerAspSerSerAlaSerAlaP 210
1009 CGAACAGCTCTTCTGCTGGCGCTTCTTGATACGAATCATCATGTGTG 960
210 roAsn...LysCysTyrAlaProSerAlaIlePro..... 220
959 TAGCGTCCCAATTAATCGTGGCGGCTTGATATGCGCGCTTCATATC 910
221 ...ThrProGlnArgThrSerThrPro...GlyLeuAlaLeuPheProG 235
909 CGAACCCGAA.....ATCAGCGGCTTCTGTCTGTCA 878
235 yProProSerProValAlaAsnSerThrSerThrProLeuThrLeuPro 252
877 CCAATTCGCC....GCAGTAATTTGCATCTTTCGACCCAAA 837
252 aGlnSerProLeuAlaThrAlaAlaSerAlaSerThrSerAlaProVal 268
836 ACGGTACCCAGAGAGCGGCTTGTGACTTGAACACCACCTAGGCAAT 787
269 SerCysGlySerSerAlaSerLeuLeuArgLysProHisProGlyThr 285
786 CACGGCGCTGGTGTTCAGACGGCTGTTCACAAACAGCCGATGTAA 737
285 rAspLeuHisIleSerSerThrProAlaAlaThrThrLeuPro..... 299
736 TTAACATCTTGAATTAATGATGCTCCACACGCGTTTATTCGCCGCGC 687
300 .....ValMetIleLysThrGluProThrSerProThrPro 311
686 TCGATGAATGAATGTGCTGCCACTCAA...CCGGCAGAGATGGGGCC 640
312 Ser.....AlaPheLysGlyProSerHisSerGlyAs 322
639 GCCGAAT.....TCATGTGTTTCGATGTTCGACGATTTTCAG 602
322 nProSerHisGlyThrLeuGlyLeuSerGlyThrLeuGlyArgAlaTyrT 339
601 ACGCAGCTGTGGCGCAGCTGCTTACAAACATGATTTGCGTTCGCTC 552
339 hrSerThrSerValProIleSerLeuSerAlaCysLeuAsnProAlaLeu 355
551 AAAGCGCTCAATACCAACAGCGCGCTTGA...TCTTCGGCGGCTTC 505
356 SerGlyLeuSerSerSerSerThrProLeuAsnGlySerAsnProLeu 372
504 TTTGATATGACCGTAGGTGGAGCCAGCCGATGTCATCATGGCAT 455
372 rSerIleSerLeuProProHisGlySerSerThrProIleAlaProValP 389
454 TGACGAAGATGGCAGACGGCTCGCATCGACGAGAGAAATTTGCTGAC 405
389 heThrAlaLeuProSerPheThrSerLeuThrAsnAsnProPheProLeu 405

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404 GAGCG....GTGGCAGCGCAGTCCACAAACCGGATTGATC..... 366
406 GlyAsnProSerLeuAsnProSerValSerLeuProGlySerLeuLeuAl 422
365 .....AGGTTGGCGGCGCACTTCTTCGCCCTTAAGTTTGGCAGGCGTT 323
422 atThrSerSerThrAlaAlaThrSerThrSerLeuProHisProSerSerT 439
322 CAGTGCGGTAGCGTTCAAACTCGATTGCTGTTGCCCTTCACGCGCAATC 273
439 hrAlaAlaValIleuSerGlyLeuSerAlaSerAlaProValSerAlaAla 455
272 ACGACGACTGAGTAGTACGGCTTTTGCCACGGTGAATGAGCGGCGATT 223
456 Pro.....PheProLeuAsnLeuSerThrAlaVala 465
222 GCCTGAAGCGCGCGCAGTAACACACACGCGGATCTTGTGCTCA 173
465 lProSerLeuPheSerVal.....ThrGlnGlyProLeuSerSerSerA 480
172 ACAGCACTTGCGCTTTTTCAGCGCATC....CCTTCCTTGACTTTC 129
480 snLeuSerTyrProGlyPheSerValSerAsnThrProSerValThrPro 496
128 ATGAGAGGCGCATACCGCATATCTTCGCCAAGCAGCGCATTCGCT 79
497 AlAlaLeuProSerPheProGlyLeuGlnAlaProSerThrValAlaAlaVala 513
513 lThrProLeuProValAlaAlaAlaThrAlaProSerProAlaProValLeuP 530
31 GCAGTTT 24
530 roGlyPhe 532

seq.name: /SIDSL/ycdata/geneseq/geneseq-emb1/AA2001.DAT:ABB64198
seq_documentation_block:
ID ABB64198 standard; Protein: 2406 AA.
XX
AC ABB64198;
XX
DT 26-MAR-2002 (first entry)
XX
DE Drosophila melanogaster polypeptide SEQ ID NO 19386.
XX
KW Drosophila; developmental biology; cell signalling; insecticide;
XX pharmaceutical.
XX Drosophila melanogaster.
XX
PN WO200171042-A2.
XX
PD 27-SEP-2001.
XX
PF 23-MAR-2001; 2001WO-US09231.
XX
PR 23-MAR-2000; 2000US-191637P.
XX
ER 11-JUL-2000; 2000US-0614150.
XX
PA (PEKE ) PE CORP NY.
XX
PI Venter JC, Adams M, Li PWD, Myers EW,
XX
DR WPI; 2001-656860/75.
XX
DR N-PSDB; ABL0830L.
XX
PT New isolated nucleic acid detection reagent for detecting 1000 or more
XX genes from Drosophila and for elucidating cell signalling and cell-cell
XX interactions -
XX
PS Disclosure; SEQ ID NO 19386; 21pp + Sequence Listing; English.

```

XX The invention relates to an isolated nucleic acid detection reagent capable of detecting 1000 or more genes from *Drosophila*. The invention is useful in developmental biology and in elucidating cell signaling and cell-cell interactions in higher eukaryotes for the development of insecticides, therapeutics and pharmaceutical drugs. The invention discloses genomic DNA sequences (AB16176-AB130511), expressed DNA sequences (AB161840-AB16175) and the encoded proteins (AB161737-AB161702).

CC The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at [ftp.wipo.int/pub/published_pct_sequences](http://wipo.int/pub/published_pct_sequences).

XX Sequence 2406 AA:

Alignment_scores:
Quality: 119.50 Length: 492
Ratio: 0.511 Gaps: 27
Percent Similarity: 47.561 Percent Identity: 21.951

Alignment_block:
US-09-303-518D-125/rev x ABB64198

Align seg 1/1 to: ABB64198 from: 1 to: 2406

1340 CCTTCCTTCTCAATGTTTCCAGCATTTCGCGCAACGCGGCGGTATTC 1291
||| :
1278 ProValLeuProValAlaThrProAsnLeuSerAsnLeuProThrG1 1294
1290 GAATTTGCCGGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1241
:
1294 HisArgSerSerAspSerArg..AsnSerArgGluSerProAlaSerL 1310
1240 ATTCCAGCAACCCCAATGCTGCGGCTGTGCGATCG.....CCGACG 1197
:
1310 euLysSerThrProSerAsnIleGlyLeuAsnValSerMetAlaProThr 1326
1196 ATTAATG.....CGCAAAAGCAGGGTGGGAGAGATATCCAAAGCGAT 1153
:
1327 LeuArgSerIleThrProLeuAsnAsnSerSerAlaIleSerSerGlyAl 1343
1152 CAGCGCTCGTAAATGCAATGCGCAGCAGCAGCGGCTGCGCGCGTTGA 1103
1343 AserGln.....ProValSerValValProSerAlaAsnSerT 1357
1102 CCGCTGTGTACTGAAAGACTTTGTTTTCAGAGAAATGCCCGAGGTT 1053
||| :
1357 hVala...LeuSerMetSerAsn.....ProHisIle 1366
1052 GThA.....CGCGTGAATGAGTATTTGTCGCGC..... 1026
1367 SerHisSerHisValProAlaThrAlaSerGlyAlaPheSerSerSe 1383
1025TCGCGCGCAACCCAGCCAGCAAGCTTTG.....CTGCGGC 989
:
1383 rAlaAlaIaGlyThrSerThrProAsnSerGlyLeuSerThrLeuAlaV 1400
988 CTTCCTTCGATTAACGGAATCTGATTTGTGTACGTCGCCAAATACGTCGC 939
:
1400 aLThrSerLeuSerThr.....SerAla 1407
938 GCGCCTGTGTAATGCGCGCTTCAATACGAGCAAGCAATCAGCGGTT 889
||| :
1408 AlaPro.....GlnProHisSerHisPheProGlnSerThrGlnMe 1421
888 G.....TCGTGTG..... 882
1421 tLeuProGlnSerGlyAsnPheSerSerValSerHisLeuThrThrN 1438
881TCACCAATTCGCGCGAGTATTTGGATATTCGATACCTTCGCA 843
1438 tSPrometSerSerGlnAsnGlnPrometValaLargCysGlySer..... 1452

842 CCCAAAGGTACGACAGAGCGCGTTTGTGACTTGAAACGACTAG 793
||| :
1453ThrLeuLysSerIleSerIleSerAlaAlaIleThrAlaProProSe 1467
792 GGCATACAGCGCGCTCGTGTTCAGACGGCTTTCGCAACAAACGCGCAA 743
||| :
1467 rAlaAlaAla.....AlaValSerAsnPheThrProS 1478
742 TGGTAAATACATCTTGATATTAATGATGTCACACAGGTTTATTCGCGCG 693
||| :
1478 eValLeu..... 1480
692 ACCGCTCGATGAATGATGCGTCCACTCAACCGCAGATGCGG 643
1480 1480
642 GCGCGCAATTCATGTGTTTCGATGTTGGCAGCATTTTCAGACGCGACT 593
:
1481AlaValGlnSerLeuThrThrAlaValThrSerSerSers 1494
592 CTGCGCGCAGCTGCTTACAAACATGATTTTGCCTTCGCTC..... 552
||| :
1494 eSerProSerThrLeuSerSerSerValIleGlnLysValIleSerPro 1510
551AAACGCTCAATACCAACAGCGCGCTTGAATTCCTCGGC 511
||| :
1511 LysGlnGluSerProCysAsnLysAspArgAspSerSerTyrSerSerTr 1527
510 GCGCTCTTGTGATATACGCTAGGCTGGCGAGCGCAGGATGTGTGTCCA 461
||| :
1527 oAlaAsnAlaValValThrCysAlaProThrThrProIleValSer. 1543
460 TCCGATTGACAGATGGCGAAGCTCGGCAATCGACGCGAGGAAATTTTG 411
:
1544SerGlySerAlaArgProThrProProLeu 1553
410 CTGAAC.....GGACGGGTGCGCAGCGCAGCTCCACA 379
||| :
1554 SerAsnCysThrSerMetGlyIleGlyMetValAsnAlaAla..SerThra 1570
378 ACCGATGTGATGATGAGCTGCGCGCAGCTTGTTCGCGCTTAAGTTGCCA 329
||| :
1570 LArg.....SerSerCysAsnAlaIle..SerProLeuSerIlePro 1584
328 GCGCTTCAGT.....GCGTAGCGTTCGAAC 303
1584 tArThrAlaGlyIleHisValSerAlaThrAsnProSerPheGlnSerSer 1600
302 TCGATTTGTCGTGCT..... 285
1601 SerTyrPheProThrProLeuAlaProProProSerSerProSerProAl 1617
284TCAACGGCATCAACGACTGACGTAAGACGCTTTTCGCA 243
1617 aThrSerSerAlaAlaIleIleSerSerSerAlaSerGlnPheAsnPro 1634
242CGGTGAATGCGCGGAGTTTGGCTGAGGCGCGGAGTAAC 201
:
1634 lAlaValSerHisSerMetSerSerIleValIThrThrAlaGlyAlaThrThr 1650
200 ACCACGCGCGGATTTCTTTTGTCTTCAACAGCACTTGGCTTTTGTGAC 151
||| :
1651 ThrThr.....AlaSerSerValThr..... 1657
150 GGCATCGCCTTCCTTGAATTCATCGAGGGCGGATACCGCATATTTCTT 101
:
1658GlnProSerValAlaAlaIleSerAsnProValIThrAsnThrProH 1673
100 GCGCAAGCAACGCACTTCGGTA 78
||| :
1673 tSPrometSerSerAlaGluSerLeu 1680


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967 .AsnAsnAspLeuGlyAlaGlyMetAlaValAlaMetLysAspLeuGluM 983
809 TCACAAACCGCCGCTTGGCTACCGCTTTGGTGGCAAGTATCGCA 858
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
983 et.....ArgGlyAlaGlyAsnValLeuGlyAlaGlyLysSerGly 996
859 ATTACTGCG.....GGCGAATT 875
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
997 HisLeuAlaGlyValGlyPheAspLeuTyrValArgLeuValGlyGluAl 1013
876 GGTTCACACAGCAACCGCGTATTCGCTTCGCTTCGCTTCGCTTCGCTTC 925
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1013 aValGluValArgAlaLeuAlaAspGlyLysValValAspGlyThrV 1030
926 TT.....ACACAAAGCGCGCAC... 942
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1030 alTyGlyProLysGluLeuArgValAspLeuProValAspAlaHisLe 1046
943 .....GATTATTGGACGCGTACCAACATCATGATTCCGTTATCGAAGA 986
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1047 ProGluLysTyrLeAsnAlaGluArgLeuArgLeuGluLeuTyrArgL 1063
987 AGCCCGCAGCAAGAGCTGTTCGCTGCGCTGCGCGCGCAGCGCAAAAT 1036
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1063 yLeuAlaLeuLysSerGluSerGluValAspLeuArgLeuAlaValGlu 1079
1037 ACTCCATCAGCGGTACACCCCTCGCGCATTTCTTCAAAAACAACCTTTC 1086
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1079 uMetGluAspArgTyrGlyProLeProGluGluValGluArgLeuVal 1096
1087 AGTTCACACAGCGCGTCAACGCGCGCGCGCAT..... 1124
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1096 lValSerArgLeuArgHisLeuMetArgGluAlaHisLeuThrAspLe 1112
1125 GGTCCGATGTGCTTACGAGCGCGTATGCCCTGATATCTGCCCA 1174
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1113 AlaValGlnGlyThrArgIleLysValHisProValAspLeuAlaAsp 1129
1175 CC.....CTGCTTTGGCGGATTTAATGCTGCGATACGACAGCGCG 1218
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1129 eArgLysValArgLeuLysArgLeuPheProGlyAlaThrTyrArgAla 1145
1219 CAGGCATTTGGTGTCTTGAATTTGACGAGAAAGACCTGCTTGTGCAG 1268
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1146 ..AlaAlaLysAlaLeuGlnLeu.....Se 1153
1269 CTTCGCTGCGCGGCAAAATACGAATACGCGCGCTGTTGCCCAAGTG 1317
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1153 rPheProLysThrGlyAsnLysValThrAspProLeuLysArgAspVal 1170
1318 .....CTGGAA 1323
1170 sPLeuLeuGlnTyrPValAlaAsnPheLLeSerAsnMetPheAsnLeuGln 1186
1324 ACCATTGAGAAAGAGGC 1341
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1187 GluLeuAspValArgGly 1192

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seq_name: /SIDSI/gcgsdata/geneseq/geneseq_emb1/AA2001.DAT.AAB59827

seq_documentation_block:

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ID AAB59827 standard; Protein; 1592 AA.
XX
AC AAB59827;
XX
DT 04-APR-2001 (first entry)
XX
DE Protein #4 encoded by TutsD/E gene.
XX
KW Toluene degradation; enzyme; waste degradation; Tuts; TutsD.
XX
OS Thauera aromatica.
OS Xanthomonas maltophilia.

```

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OS Geobacter metallireducens.
OS Azobarcus toluyticus.
XX
XX WO200072650-A2.
XX
XX 07-DEC-2000.
XX
XX 24-MAY-2000; 2000WO-US14298.
XX
XX 01-JUN-1999; 99US-0323872.
XX
XX (UYOH-) UNIV OHIO.
XX
XX Coschigano PW;
XX
XX WPT; 2001-041080/05.
XX
XX N-PSDB; AAF23627.
XX
XX Composition comprising toluene degrading enzyme useful for biological
XX treatment of organic compounds, especially for degrading toluene or its
XX analogs.
XX
XX Disclosure; Fig 12; 122pp; English.
XX
XX The present invention relates to toluene degrading enzyme genes and
XX proteins tuts (see AAF23629 and AAB59831), tuts (AAF23630 and AAB59832),
XX tuts (AAF23631 and AAB59833) and tuts (AAF23632 and AAB59834). The
XX toluene degrading enzymes are homologues of pyruvate formate lyase. The
XX toluene degrading enzymes are useful for biological treatment of organic
XX compounds and in particular for the degradation of toluene and its
XX analogs contained in liquid or solid waste source. The present sequence
XX is a protein sequence encoded by toluene degrading enzyme gene, TutsD/E.
XX
XX Sequence 1592 AA:

```

```

alignment_scores:
  quality: 118.00      length: 584
  ratio: 0.599        gaps: 34
  percent similarity: 33.733  percent identity: 22.774

```

alignment_block:

US-09-303-518D-125/rev x AAB59827 ..

Align seg 1/1 to: AAB59827 from: 1 to: 1592

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1312 TGCGCAACAGCGGCGGCTATTCGATTTGGCCGCGCA..... 1274
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
707 CysThrSerArgGlyArgSerArgCysSerProAspArgCysArgT 723
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1273 .CGAAGCTGCACAAAGCGAGGCTCTTCGTCAATTCACAGCAACCA. 1226
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
723 pSerArgCysSerSerProSerProArgArgCysProProSerProA 740
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1225 .....ATGCCCTGCGCGCTGTGCGTATCGCGCAGCATTAATCGCGCAAAA 1181
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
740 lAglyAlaProGlyAlaThrCysSerArgArgProPheSerArgSer 756
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1180 GCAGGGTGCGCAGCATATCCAAAGGCATCGCGCTCGTAGTACCAATC 1131
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
757 AspSer...AlaGly..ProArgAlaAlaArgPheArgArgCysArgAs 771
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1130 GGCACCATGCGCGCGCTGCGCGCGCTTGACGGCTGTGTTGAACCTGAGAG 1081
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771 pAlaCysGluArgArgAlaArg..... 778
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1080 TTTGTTTTCAGAAATGCGCAGAGGTGTACGCTGATGAGATTTGT 1031
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
779 .....Cys 779
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1030 CCG.....GC 1026
   ||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
780 ProGlyProArgSerArgProSerLLeArgArgGlySerArgAspArgSe 796

```

```

1025 TCGGGCGCAACCCAGCCGACGCTCTTTGCTGCGGCTTCTTGATAC 976
      : ||||| :|||
796 rArgAlaSerArgSerArgGlySerProLeuGlySly..... 808
975 GGAATTCGATGTGTGTAGCTCCCAATATCGTGGCGGCTGTGTAA 926
      : :||| :|||
809 .....AlaThrAlaThrSerCysProAlaArgArg..... 818
925 TCGCGGCTTCATATACCGAAGCAATACGGGCTGTGTGTGCAAC 876
      ||| |||||
819 ...ArgCysSerIle.....GlyAlaSerSerGlyCysProHisProPr 832
875 AATTGCGCGGAGTAATTTGGCATCTTTCGACCCCAAAACGTAACGAA 826
      :|||
832 ovalArg..... 835
825 GAGGGGGGTTTGTGACTTGAGAACCCATAGGCAATACGCGCTCGG 776
      :|||
835 rGserProVal.....AsnSerSerIlySerArgAlaHisArgArg 847
775 TGTTCAGAGGCGCTGTGCAAAACAGCGCATGTATATCATCTTGA 726
      |||||
848 CysThr..... 849
725 TAATTGATGTCACACGCTTTATTCGCGCGACGCGCTGATGAATG 676
      :|||
850 .....AlaArgArgGlyArgPheArgGlyPro..... 858
675 AATGTGCGTGCACCTCAACCGGAGATGAGGCGCGCCCAATCATGTG 626
      :|||
859 .....ThrSerArgAspThrGlyArgArg..... 866
625 TTTGAGTGTGGCAGCATTTTCACAGCGCACGTCGCGC..... 587
      |||||
867 ...ArgCysTrpArgTrpProArgProAlaArgCysArgCysSerArgAr 882
586 ...CAGCTGCTTACAACATGATTTTGCCTTCG..... 554
882 GTTPGLYArgProLeuTrpAlaSerGlyCysProArgAlaArgTrpArg 899
553 .....TCMAACGGCTCAATACCAACAGCGCGTTCGTAATCCTCGGCG 509
      |||||
899 rGlySerAsnTrpSerSerGlyArgSerSerAlaAlaSerProGlySarg 915
508 CTCTCTTGATTAATGACCGTAGGCTCGGCAG..... 479
916 ThrCys.....GlyArgArgValArgSerAspThrSerAl 927
478 .....CCAGCGGATTCG.....T 466
927 aArgArgSerArgCysProAlaSerSerProIleArgTrpThrGlyArgC 944
465 GTCCATCGCATTTGACGAGATGCGCA..... 440
944 ys.....ArgArgTrpArgArgProLeuGlySerSerProArg 956
439 .....ACGGCTCGGACATGACGCGCAGAAATTTGCTGAACGACGCG 399
957 AlaThrCysThrAlaArg.....CysGlyArgAspGly 967
398 GTGCGCGACG..... 389
967 yCysSerAlaPhePheGlyAsnProLeuHisArgSerLeuArgGlyProT 984
388 .....CAGTCCACAACCGG..... 374
984 rPAlaAlaProPheArgAlaHisArgSerArgSerThrThrArgArgCys 1000
374 ..... 374
1001 AlaValArgGlySerSerArgHisAspArgThrAlaSerThrArgArgPr 1017

```

```

373 .....ATTGATCAAGTTGGCGCG 355
1017 OHsLysProProLysGlyCysAlaThrAspIleHisSerGlyArgTyrC 1034
354 CACTTCTTCGCGCTTAAGTTTCCAGCGGCTTCAGGTGGTACGCT... 308
1034 yStrProAlaGlyThrAlaSerSerArgAlaAlaSerGlyAlaSerAlaLys 1050
307 CAACCTCGATTTGTCGCT.....TGCCTTCAACGGCAATACGACGTGAC 264
1051 ArgThrArgLeuArgArgArgSerCysProValArgSerProAlaArgArg 1067
263 TGAAGTACGCGCTTTTCGCGCAC.....GCTGATGCGCGCATTTT 223
1067 gGlyThrArgAlaAlaThrPheHisSerAlaCysGlySerSerArg.... 1082
222 GCCTGAAGCGCGCGCAGTAACACCA.....CGCCGCGAT 188
1083 ...ArgProSerSerGlyArgProTrpSerValProIleArgProSer 1097
187 TCTTTTGTCTTCAACACGACTTGCGCTT..... 158
1098 SerIleCys...GlyArgAlaValGlyLeuThrSerProSerSerProle 1113
157 .....TTTGACGGCATCGGCTTCCTGACTTTCATTCAGAGGCGG 118
1113 uAsnArgProPheAlaArgArgSerAlaPro.....AlaSerThrProC 1128
117 CATACCGGCAATTTCTTCCCAACAGCACGCGACTTCGTAATGCGCGGC 68
1128 yArgArgHisAsnArgAlaArgArgTyrGlySerArgArg.....Prohe 1142
67 CGTCGTAA..... 59
1143 ArgArgArgPheAlaCysSerTrpSerSerGlnHisAspProAlaSerG 1159
58 .....CGGCTTGCTCCGCTGCGCGCGAGATGGG 31
1159 nAspProGlnArgGlyThrCysProLeuArgAsnAlaCysProGlyTrpA 1176
30 CA 29
1176 Ia 1176
seq_name: /SIDSL/gcgdata/geneseq/geneseqp-emb1/AA2001.DAT:ABG22380
seq_documentation_block:
ID ABG22380 standard; Protein; 2447 AA.
XX
AC ABG22380;
XX
DT 18-FEB-2002 (first entry)
XX
DE Novel human diagnostic protein #22371.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
XX food supplement; medical imaging; diagnostic; genetic disorder.
XX OS Homo sapiens.
XX PN WO200175067-A2.
XX PD 11-OCT-2001.
XX PF 30-MAR-2001; 2001WO-US08631.
XX PR 31-MAR-2000; 2000US-0540217.
XX PR 23-AUG-2000; 2000US-0649167.
XX PA (HYSE-) HYSEQ INC.
XX PI Drmanac RT, Liu C, Tang YT;
XX

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DR WPI: 2001-639362/73.
DR N-PSDB; AAS86567.

PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity -

PS Claim 20; SEQ ID NO 52739; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful for treating
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. ABG00010-ABG30377 represent novel human
CC diagnostic amino acid sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pcl_sequences.

SO Sequence 2447 AA:

alignment_scores:
Quality: 118.00 Length: 473
Ratio: 0.527 Gaps: 23
Percent Similarity: 47.357 Percent Identity: 23.890

alignment_block:

US-09-303-518D-125 x ABG22380 ..

Align seg 1/1 to: ABG22380 from: 1 to: 2447

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18 AGCTTAAACCTGCCATCGCGGACGACGCAAGC.....CG 58
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1799 ArgSerArgSerLysSerArgLysArgSerValSerLysGluLysArg 1815
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
59 TTTACGACGCGCGCCGATTACCGAAGTCGCGTTCGCGAAGAAATAT 108
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
1815 GlySerArgSerProLysHis...ArgSerLysSerArgLysArgLysArgL 1831
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
109 GCGCGTATGCGCGCCCTCGATGAAGTCAAGAGGCGATCGCGTCAAAA 158
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
1831 yArgSerSerSerArgSerArgSerValArgLysThrValArgLysArgSerArg 1847
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
159 AGCCCAAGTCTGTTGAAGA.....CAAAAGAAATCCGG 193
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
1848 ThrProSerArgArgSerArgSerHisThrProSerArgArgArgArgSe 1864
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
194 GCGTGGGTTTACTGCGCGGCTTCAGGCAAAATCGCGCATTCACCGT 243
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
1864 rArgSerVal.....GlyArgArgArgSerPheSerIleSerPro 1878
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
244 GCGCAAAAGCGCGTACT..... 260
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
1878 eArgArgSerArgThrProSerArgArgSerArgThrProSerArgArg 1894
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
261 .....TCAGTCAGTCGTGATTCGCGTGAAGCAACGCAAAATCGAGT 304
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
1895 SerArgThrProSerArgArgSerArgThrProSerArgArgSerArgThr 1911

```

```

305 TTGAACGCTAACGCACCTGAAGCGCTGGCAACTTAAGCGCGGACGAAGTGC 354
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
1911 rProSerArgArgSerArgThrProSerArgArgArgSerArgSerArg 1928
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
355 GCGCGCAACCT.....GATCCAAATCCGGTTG..... 381
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
1928 aValArgArgArgSerPheSerIleSerProValArg...LeuArgArgSe 1944
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
382 .TGACGTGCGCTGCGACCGCGCTTCAGCAAAATTCCT..... 420
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
1944 rArgThrProLeuArgArgArg...PheSerArgSerProIleArgArgL 1960
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
420 ..... 420
1960 yArgSerArgSerSerGluArgGlyArgSerProLysArgLeuThrAsp 1976
   .....GCGGTGATGCGCGAGCGGTTCGCCAT 446
421 ..... 421
1977 LeuAspLysAlaGlnLeuLeuGluIleAlaLysAlaAsnAlaAlaLys 1993
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
447 CTTCGTCATATGC.....ATGACACCAATCCGCTGCGTCCGAC 487
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
1993 tCysAlaLysAlaGlyAlaProLeuProProAsnLeuLysProAlaPro 2010
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
488 CTACGTCATATTCAAAGAACCGCGAGATTCAAACCGCGCTTTG 537
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
2010 rProThrIleGluGluLysValAlaLys.....LysSerGlyGlyAla 2024
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
538 GTATTCAGCGGTTTCAGCCGCAACCAATCCATGTTGTAGGCACTGG 587
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
2025 ThrIleGluLeuLeuThrGluLysCysLysGlnIleAlaGlnSerLysG 2041
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
588 CGCAGCGTCGCGCTCTGAAATGCTGCCAATCGCAATCGAATCAATTCG 637
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
2041 uAspAspAspValIleValAsnLysProHisValSerAspGluGluG 2058
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
638 GCGGCGCG.....CATCTCCGCGTTTGAATGCGACGAC... 672
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
2058 LuGluProProPheThrHisHisProPheLysLeuSerGluProLysPro 2074
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
673 ATTGATTCATCGAGCGCGGTCGCAATGAAACCGTGTGACATCA 722
   |||||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
2075 IlePhePheAsnLeuAsnIleAlaAlaLysProThr.....Pr 2088
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
723 TTATCAAGATGTAATTCATTCATGCGCGCTTGTTCGAACGCGCTGTA 772
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
2088 oProLysSerIleValIleThrLeuThrLysGluPheProValSer..... 2102
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
773 ACACCGAGCGCGTATTCGCTAGGTGCTTCAGTCAACCAACCGCGC 822
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
2103 .....SerGlySerGln...HisArgLysLys 2110
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
823 CTCCTTGCCTACCGTTTCGGTGCAGAAAGTATTCGCAAAATTCGCGGCGA 872
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
2111 GluAlaAspSerValTyrGluGluThrAlaProValGluLysAsnGlyG 2127
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
873 ATTGGTTGACACAGACAAACCGCGTATTCGCTTCGATTCGAACCGCGC 922
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
2127 uGluAsnLysAspAspAspValPheSerSerAsnLeu..... 2140
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
923 CGATTACAAAGCGCGCGACGATTAATTTGGAGCCTTACACATCAATTA 972
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
2141 ..ProSerGluGlyArgValLysArgGlnGlyAlaArgValArgGlnMet 2156
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
973 .....TCGTTATGCAAGAGCGCGACGCAAGAGAGCTGT 1007
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
2157 LysGlnProAlaAlaSerHisLeuThrValThrArgCysAsnSerLeu 2173
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
1008 GCGCTGGGTTGCGCGACCGCGCAAAATCTCCATCAACGCGTCAACACC 1057
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
2173 sely...ThrLysProGlnSerGluLysHisArgIleAlaGluAsnSer 2189
   :|||:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|:|
1058 TCGGCCATTTCCTGAAGAAC.....AAACTCTTCAGATTC 1092

```

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      2189  a l l e t h s e r l e u p r o a s n i l e g l y p r o s e r l e u h i s l e u t r e u l a 2205
      |||
      2206  a s n l e u s e r g l y g l y l e u s p a r g a r g l y s n i l l e u a r g p h e a s 2222
      |||
      2222  n t h r a r g t h r g l y l e u g l u y s v a l l i e p r o v a l a s p l e s e r t h r a l a m 2239
      |||
      1178  t g c t t t t g c g c a t t t a 1194
      |||
      2239  e t s e r g l u a r g a l a l e u 2244

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seq_documentation_block:
ID   AAB59817 standard; Protein; 999 AA.
AC   AAB59817;
DT   04-APR-2001 (first entry)
DE   TultD protein #8.
KW   Toluene degradation; enzyme; waste degradation; TultD.
OS   Thauera aromatica.
OS   Xanthomonas maltophilia.
OS   Geobacter metallireducens.
OS   Azotarcus toluilyticus.
PN   WO200072650-A2.
PD   07-DEC-2000.
PF   24-MAY-2000; 2000WO-US14298.
PR   01-JUN-1999; 99US-0323872.
PX   (UYOH-) UNIV OHIO.
PA
XX
PI   Coschigano PW;
DR   WPI, 2001-041080/05.
DR   N-PSDB; AAF23625, AAF23627.
PT   Composition comprising toluene degrading enzyme useful for biological
PT   treatment of organic compounds, especially for degrading toluene or its
PT   analogs
XX
PS   Disclosure: Fig 5; 122pp; English.
XX
CC   The present invention relates to toluene degrading enzyme genes and
CC   proteins tult (see AAF23629 and AAB59831), tult1 (AAF23630 and AAB59832),
CC   tult2 (AAF23631 and AAB59833) and tult3 (AAF23632 and AAB59834). The
CC   toluene degrading enzymes are homologues of pyruvate formate lyase. The
CC   toluene degrading enzymes are useful for biological treatment of organic
CC   compounds and in particular for the degradation of toluene and its
CC   analogs contained in liquid or solid waste source. The present sequence
CC   is a protein sequence for toluene degrading enzyme, TultD.
XX
SQ   Sequence 999 AA:

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alignment_scores:
  Quality: 117.00      Length: 570
  Ratio: 0.582         Gaps: 33
  Percent Similarity: 35.263  Percent Identity: 23.158

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alignment_block:
US-09-303-518D-125/rev x AAB59817

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112  CysThrSerArgGlyArgSerArgCysSerProSerArgCysArgTrp 128
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1273  .CGAAGCTGCACAAGCGAGGTCTTTCGTCGAATTCACCAACCCA 1226
|||
128  pSerArgCysSerSerProSerProArgArgCysProProSerSerProA 145
|||
1225  ....ATGCCGTGCGCGCTGCGGTATCGCGGAGCAATTAATGCGGAAA 1181
|||
145  IacIyAlaProGlyAlaThrCysSerArgArgProPheSerArgSerArg 161
|||
1180  GCAGGTGGCGAGATATCCAAAGGCATCAGCGCTGTGAATACCAATC 1131
|||
162  AspSer...AlaGly...ProArgAlaAlaArgPheArgArgCysArgAs 176
|||
1130  GGCACCATGGCGCGCGCGCGCTGACGCGCTGTGTGAATTAAGAG 1081
|||
176  pAlaCysGluArgArgAlaArg..... 183
|||
1080  TTTGTTTTCAGGAATATGCGCGAGGTTGTACGCGTGAATGATTTGT 1031
|||
184  ..CysProGlyProArgSer.....Ala 190
|||
1030  CCGGCTGCGCGCGCAACCCAGCCGAACGCTCTTTCGCGGCTTTCG 981
|||
191  ProSerIleArgArgGlySerArgAspArgSerArgAlaSerArgSerArg 207
|||
980  ATACGGAATCTGAT...TGTGTAGCGGTCCCAATATATATCGTGGCC 934
|||
207  gSerArgGlySerProLeuCysGlyAlaThrAlaThrSerArgProArg. 223
|||
933  TTGTGTAATCGCGCGCTTCATACCGCAACCGGAATACGCGGTGTCTG 884
|||
224  ....ArgAlaArgCysSerIleGlyAlaSerSerGlyCysPro 236
|||
883  TGTCAACCAATTCGCCCGCAGTAATTTGCGATCTTTCGCCACCAAAAG 834
|||
237  HisProProValArg..... 241
|||
833  GTACGCAAGAGCGCGGTTTGTGACTGAGAACACCACTAGGCAATCAC 784
|||
242  ....ArgSerProVal.....AsnSerSerIysArgAlaH 252
|||
783  GCGCTCGGTGTTTCAGACGCGCTGTTCAAACAAAGCGCAATGTTAATTA 734
|||
252  IsArgArgCysThr..... 256
|||
733  CATCTGATTAATGATGTGCCAAGCGGTTTATTCGCCGCGACCGCTCG 684
|||
257  ....AlaArgArgGlyArgPheArgGlyPro..... 265
|||
683  ATGAATGATATGTGCTGCCACTCAACCGCGAGATGCGCGCGCGAA 634
|||
266  ....ThrSerArgAspPheCylArgArg. 273
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633  TTCAATGTTCGATGTTGCGACGATTTTTCAGACGCGACGCTGCGC. 587
|||
274  ....ArgCysTrpArgTrpProArgProArgArgCysArgCys 286
|||
586  ....CAGTCCCTTACAAACATGAGATTTTTCGCTGCG..... 554
|||
287  SerArgArgTrpGlyArgProLeuTrpAlaSerArgCysProArgAlaAr 303
|||
553  ....TCAACGCGCTCAATATACCAACAGCGCGGTTTGAATATC 517
|||
303  gTrpArgArgGlySerAsnTrpSerSerArgGlyArgSerSerAlaAlaSer 320
|||
516  CTCGCGCGCTTCTTGTGATTAATGACCGTAGGCTCGCGAG..... 479
|||

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320 rOLysArgThrCys.....GLyArgValArgSerAsp 331
478 .....CCAGCGATTGG.. 467
332 ThrSerAlaArgSerArgCysProAlaSerSerProIleArgTrp 348
466 .....TGTCATCGCATTCAGCAAGATGGCGA..... 440
348 rGLyArgCys.....ArgArgTrpArgArgProLeuGLyCys 361
439 .....ACGGCTGGCATTCAGCGAGAGATTTCGTGA 407
361 eRPtoAlaThrCysThrAlaArg.....CysGLy 371
406 ACGGACGGGTCGCGACG..... 389
372 ArgAspGLyCysSerAlaRheRheGLyAsnProLeuHisArgSerLeuArg 388
368 .....CAGTCCACCAACC 376
388 gGLyPRoTrpAlaAlaProRheArgAlaHisArgSerArgSerThr 405
375 GG..... 374
405 rGArgCysAlaValArgGLySerSerArgHisAspArgThrAlaSerThr 421
373 .....ATTGATCAGG 363
422 ArgArgProHisLysProProLysGLyCysAlaThrAspIleHisSerGI 438
362 TTGCGCGCGCATCTTCGCGCGTTAAGTTTGCAGCGCTTCAGTGGCTA 313
438 YArgTrpCysTrpProArgThrAlaSerSerArgAlaAlaSerGLyAla 455
312 GGGT...CAACTGCATTTCGTCT...TGCTTCAACGGCATCA 272
455 eRAlaLysArgThrArgLeuArgArgArgSerCysProValArgSerPro 471
271 CGACTGACTGAAGTAACGCGCTTTGCGCAC.....GGTGAATCGCG 231
472 ArgArgArgGLyThrArgAlaAlaTrpHisSerAlaCysGLySerSerSe 488
230 GGGATTTCCTCCGAGCGCGCGCAGTAACACCA.....C 196
488 rArg.....ArgProSerSerGLyArgProTrpSerValProIleA 502
195 GCCCGGATTCTTTGCTTCAACAGCAGCATTGGCCTT..... 158
502 rGProSerSerIleCys...GLyArgAlaValGLyLeuHisSerProSer 517
157 .....TTTGAGCGGCAATCGCTTCCTTGACTTTGATC 126
518 SerProLeuAsnArgProRheAlaArgArgSerAlaPro.....AlaSe 532
125 GAGGCGCGCATACCGCATATCTTCGCCAAGCAACGCGACTTCGGTAAT 76
532 rTrpProCysArgArgHisAsnArgArgArgTrpGLySerAlaArg... 547
75 GCGCGGCGCGTCGTAA..... 59
548 ...ProRheArgArgArgRheAlaCysSerTrpSerSerIleHisAspPro 563
58 .....CGGCTTGCTCGCGGTGCGCC 39
564 AlASerGIAspProGIAsnArgGLyThrCysProLeuArgAsnAlaCysSer 580
38 GCGATGGGCA 29
580 OGlyTrpAla 583

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seq_name: /SID1/gcadata/geneseq/geneseq-emb1/AA2001.DAT.AAB27242

seq_documentation_block:

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ID AAB27242 standard; Protein: 571 AA.
XX AC AAB27242;
AC 27-MAR-2001 (first entry)
DT 27-MAR-2001 (first entry)
XX DE Human EXMAD-20 SEQ ID NO: 20.
XX DE Human EXMAD-20 SEQ ID NO: 20.
XX KW Extracellular matrix and adhesion-associated protein; EXMAD; cancer;
XX KW Inflammation; reproductive disorder; cardiovascular disorder;
XX KW Immune disorder; musculoskeletal disorder; developmental disorder;
XX KW gastrointestinal disorder; cell proliferation disorder.
XX OS Homo sapiens.
XX PN WO200068380-A2.
XX PD 16-NOV-2000.
XX PF 10-MAY-2000; 2000MO-US12811.
XX PR 11-MAY-1999; 99US-013643.
XX PR 23-AUG-1999; 99US-0150409.
XX PA (INCYTE) INCYTE GENOMICS INC.
XX PI Bandman O, Hillman JL, Tang YT, Lal P, Yue H, Baughn MR, Lu DM;
XX PI Azimzal Y;
XX DR WPI: 2001-007395/01.
XX DR N-PSDB; AAC66909.
XX PT Isolated polynucleotide encoding extracellular matrix or
XX PT adhesion-associated protein (EXMAD) useful for diagnosing, treating, or
XX PT preventing disorders associated with expression of EXMAD such as
XX PT proliferative, immune and genetic disorders -
XX PS Claim 1: Page 106-107; 129pp; English.
XX CC The present invention provides the protein and coding sequences for 25
XX CC novel extracellular matrix and adhesion-associated proteins (EXMADS).
XX CC These are designated EXMAD-1, EXMAD-2, EXMAD-3, EXMAD-4, EXMAD-5,
XX CC EXMAD-6, EXMAD-7, EXMAD-8, EXMAD-9, EXMAD-10, EXMAD-11, EXMAD-12,
XX CC EXMAD-13, EXMAD-14, EXMAD-15, EXMAD-16, EXMAD-17, EXMAD-18, EXMAD-19,
XX CC EXMAD-20, EXMAD-21, EXMAD-22, EXMAD-23, EXMAD-24 and EXMAD-25. They are
XX CC useful in the prevention and treatment of cancers, cell proliferation,
XX CC cardiovascular, reproductive, immune, musculoskeletal, developmental and
XX CC gastrointestinal disorders and inflammation.
XX SO Sequence 571 AA;

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alignment_scores:

Quality:	116.00	Length:	472
Ratio:	0.560	Gaps:	20
Percent Similarity:	43.856	Percent Identity:	19.703

alignment_block:

US-09-303-518D-125/rev x AAB27242

Align seg 1/1 to: AAB27242 from: 1 to: 571

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1340 CCTTCCTTCGTAAGTTCGACGACATTTCGCAACAGCGCGGTAATTC 1291
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23 PrometMetProThrThrSerGLyThrSerGIAsnSerSerPheAs 39
1290 GTATTGCCCGGCGAGACGAGCTGCACAAAGCGTCTTCGTCCA 1241
    :|||||:::|||||:::|||||:::|||||:::
39 nThAlaLysThrSerThrSerLeuHis.....SerHisThrSers 53
1240 ATTCCAAAGCAACCAATGCTGCGCGGTGTCGCGTA..... 1206
    :::::|||||:::|||||:::|||||:::
53 eRThrHisHisProGIuValThrProThrSerIleThrAsnIleThrLeu 69

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1205 .....TCGCGACGATTAAATGCGCAAAAG 1180
70 AsnProThrSerIleGlyThrThrProValAlaHisThrSerAl 86
1179 CAGGTTGGGACGAGATCCAAAGGACGCGCTGTAAGTACCAATCG 1130
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86 ThrSerSerAlaLeuThrThrProThrThrHisSerProThrG 103
1129 GCACC.....ATGCGCGGTGCGCGCTGACGGCTGCTGTGACCTG 1086
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103 LysSerSerProIleSerSerThrGlyProMetThrAlaThrSerPheGln 119
1085 AAGAGTTGTTTTCAGAAAATGCGCGAGGTGTACCGCTGATGAGTA 1036
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120 ThrThrThrTyrTyrThrPro..... 126
1035 TTGTGCGCGGTGCGCGCAACCGCAACAGCTTTGTGCGGCTT 986
127 .....ProSerHisProGlnThrThrLeu..... 134
985 CTTCGATACGGAATCTGATTGTGTAGCTCCCAATAATCGTGGCG 936
134 ..... 134
935 CCTGTGTAAATCGCGCGCTTCAATACGAA.....CCGGAATC.. 897
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135 ProThrHisValProProPheSerThrSerLeuValThrProSerThrH 151
896 .....ACGCGGTGTCTGTGTCACCAATCGC 869
151 sThrValIleIleThrThrHisThrGlnMetAlaThrSerAlaSerIleH 168
868 CCGCAGTAATTTGGATACCTTCGACCCAAAGATGACGAGAGCGC 819
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168 IsSerThrProThrGlyThrValProProThrThrLeuValAlaThr 184
818 GGTGTGTGACTGAGAACCACTAGGCAATC..... 786
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185 GlySerThrHisThrAlaProPheMetThrValThrThrSerGlyThrse 201
785 .....ACGCGCTCGGTTCAGACGCGCTTGCACCAACAAAGG..... 747
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201 rGlnThrHisSerSerPheSerThrAlaThrAlaSerSerPheIles 218
746 .....CCAAATGTAATTAATCATTTGATTA 723
218 eISeSerSerIrpSerSerIrpLeuProGlnAsnSerSer..... 232
722 TTGATGTCACACAGTTTATTCGCGCCGACCGCTCGATGAATGAAT 673
233 .....ArgProProSerSerProIleThrTh 241
672 GTGCTGCCACATCAACCGGAGATGCGGCGCGCAATTCATGTGTTT 623
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241 rGlnLeuProHisLeuSerSerAlaThrThrProValSerThrThAsnG 258
622 CGATGTGGACGATTTTCAGACGACGCTGCGCAGCTGCTTACAA 573
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258 InLeuSerSerPheSerProSerProSerAlaProSerThrValSer 274
572 ACATGATTTTGGCTCGGTCAAGGCTCAATACCAACAGCGCGGTTT 523
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275 SerTyrValProSerSerHisSerSerProGlnThrSerSerProSerVa 291
522 GAATGCTCGCGGCTTCTTTGATTAATGACGTAAGGCTGCGACGACG 473
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291 lGlyThrSerSerSerPheValSerAlaProAlaHisSer..... 304
472 GATGTGTTCATCGCATTCAGCAAGATGCGACGCTCGGCATCG... 426
305 .....ThrThrLeuSerSerGlySerHisSerSer 314

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```

425 .....ACGCGAGAAATTTGCTGACAGCAGCGGTGCG 394
315 LeuSerThrHisProThrThrAlaSerVal..... 324
393 CAGCGCAGTCCACAAACCGGATGTGATCAGTTGCGCGCAGCTGTCGC 344
325 .....SerAlaSerP 328
343 CGCTTAAGTTTGGCAGCGCTTCAAGTGTGAGCTTCAAACTGAT... 297
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328 rGluLeu...PheProSerSerProAlaAla.....SerThrThrIleArg 341
296 TCGTGTGCTTCAACGCGCAATCAAGACTGACTGAATACCGCTTTTC 247
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342 AlaThrLeuProHisThrIle.....SerSe 350
246 GCCACGCGTAATCGCGGATTTTGCT...GAAGCCGCGCAGTAACA 200
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350 rProPheThrLeuSerAlaLeuLeuProIleSerThrValThrValSerP 367
199 CCACGCGCGGATTTCTTTGCTTCAACAGCAGCTTGCGCTTTTGACG 150
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367 rThrProSerSerHisLeuAlaSer...SerThrIleAlaPhe...Pro 381
149 GCATCGCTTCTCTGACTTTCATCGAGGCGCATACCGCATATTTCTTC 100
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382 SerThrProArgThrThrAlaSerThrHisThrAlaProAlaPheSerSe 398
99 GCCACGCAACGCGACT 84
398 rGlnSerThrThrSer 403

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seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:ABG03530

seq_documentation_block:

ID ABG03530 standard; Protein; 599 AA.

XX ABG03530;

DT 13-FEB-2002 (first entry)

XX Novel human diagnostic protein #3521.

XX Human; chromosome mapping; gene mapping; gene therapy; forensic;

XX food supplement; medical imaging; diagnostic; genetic disorder.

OS Homo sapiens.

XX W0200175067-A2.

XX 11-OCT-2001.

XX 30-MAR-2001; 2001MO-US08631.

XX 31-MAR-2000; 2000US-0540217.

XX 23-AUG-2000; 2000US-0649167.

XX (HYSE-) HYSEQ INC.

XX Drmanac RT, Liu C, Tang YT;

XX WPI; 2001-639362/73.

XX N-PDB; AAS67717.

XX New isolated polynucleotide and encoded polypeptides, useful in

XX diagnostics, forensics, gene mapping, identification of mutations

XX responsible for genetic disorders or other traits and to assess

XX biodiversity -

XX Claim 20; SEQ ID No 33889; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and

XX polypeptide (II) sequences. (II) is useful as hybridisation probes,

CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. ABG0010-ABG30377 represent novel human
CC diagnostic amino acid sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 599 AA;

alignment_scores: Quality: 116.00 Length: 520
 Ratio: 0.563 Gaps: 36
Percent Similarity: 39.615 Percent Identity: 26.538

alignment_block:
US-09-303-518D-125 x ABG03530 ..

Align seg 1/1 to: ABG03530 from: 1 to: 599

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27 CCTGCCCATCGCGGAGACCGGACGCGTTTACGACGCGCGGCCA 76
   |||||
22 ProAlaProGlyCysArgArgAlaAlaProArgTyrSerProGlyPr 38
   |||||
77 TTTCGGAAGCGCGTGTGGGAGAGATA..TGCGGTATGCGCCCG 123
   |||||
38 o...ArgThAlaAlaGly***ArgArgMetTPrCysAlaSerAla 54
   |||||
124 TCGATGAAGTCAAGAGAGCGATGCGGTCAAAAAGGCCAAGTGTGT 173
   |||||
54 euAla**SerPro.....CysArgProArg...SerArgTyr 66
   |||||
174 TGAAGCAAAAAGATCCGGCGCTGCTTACTGC..GCCGGCTTCAG 220
   |||||
67 TPrArgAspAlaGlySerGlyTPrThrProHisCysProAlaSerAla 83
   |||||
221 GCAAAATCGCGGATTCACCGTGGCAAAA.....GCCGTACTTCAG 264
   |||||
83 atPrGlyAlaGlnGlnProValArgSerTPrGlyProArgAlaSerG 100
   |||||
265 TCAAGTCGTAGTCCGTTGAAGCAACGACGAATCGAGTTTGAACGCTA 314
   |||||
100 InSerHis..... 102
315 CGCACCTGAAGCGCTGGCAAACTTAAGCGGCAAGAAGTGGCGGCAAC 364
   |||||
103 .....CysProGlyGlyLeuArg.....AlaProProPr 112
   |||||
365 TGATCATTCGCGTGTGGACTGCGC.....TGCGACCGCTCGCTTC 408
   |||||
112 oglySerValArg..CysSerThrGln***AspCysSerSerValArgPr 128
   |||||
409 AGCA.....AAATTCCTCGCTGCATGC 431
   |||||
128 oAlaTPrSerArgSer***GlyAlaCys***GlnVal**ProArgCysP 145
   |||||
432 CGACCGCTGGCCATCTTCGATATGCGATCGACACCAATCCGTGGCTG 481
   |||||
145 rOCysArgThrProAlaThrIle.....TPrAlaPro..... 155
```

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482 CCGACCTACGTCATTATCAAGAACCCGCGAGATTTCAACCGCGC 531
   |||||
156 Pro.ProGlnGly.....ArgCysGlyProThrThrAlaPr 167
   |||||
532 CTGTGGTATTGAGCCGTTTGACCGCAACGCAAAATCCATGT..... 572
   |||||
167 rOglySerThrGlyProAlaGlyArgAlaSerLeuCysProArgArg 183
   |||||
573 .....TTGTAAGGACAGCTGGCGACAGCGCGCTGAAATAGTCGCA 616
   |||||
184 AlaHisLeuProGly***TPrProGlnLeuLeu.....CysAl 197
   |||||
617 ACAT.....C 621
   |||||
197 ahisProGlyAlaAlaSerLeuGlyLeuAlaCysGlnProHisArgGlyL 214
   |||||
622 GAAACACATGAATTCGCGGCGCGCATCTCGCGGTTGAGTGGACGCA 671
   |||||
214 ysglyThrProIleGlnGlyProAla...CysGlyThr**GlyGlyArg 229
   |||||
672 CATTCATTTCATCGAGCGCGGCGCGGCAATTAACCGTGTGACCATCA 721
   |||||
230 ArgGlySerGlyCysProGlyArg.....ProHisTh 240
   |||||
722 ATTATCAAGATGTATTATACCATTCGCGCTTGTTCACACAG..... 764
   |||||
240 rArgArgArgCys***Pro.....ProAlaProCysGlyArgArgSerA 255
   |||||
765 ..CCGCTGGAACACCGAGCGCGTATGCGCTAGAGGTCTCAAGTCAA 812
   |||||
255 laGlySerAlaHisProAlaThrProTPrProHisGlyProGlyGly 271
   |||||
813 CAACCGCGCGCT...CTTGGCTACCGT.....TTTGGTGCAGAAATAT 853
   |||||
272 GlnArgAspProGlyProAlaTyrArgGlyGlyGlnGlyAlaArgSerPr 288
   |||||
854 CGCAATTTACTGCGCGCAATGTGTTGACACAGACACCGCGGTATTC 903
   |||||
288 oAlaSerProSerGlyArg.....A 295
   |||||
904 GGTTCGATTTGAACGCGCGCATTTACAAAGCGCGACGATTTATTGG 953
   |||||
295 rGlnProAlaSerArgAlaGlyArgSerArgAlaAlaArg.....Gly 309
   |||||
954 ACGTACCAATCAATGATTCGTTATCGAAGAGCGCGACGCAAAAGC 1003
   |||||
310 ThrProGlyArgProGlu.....ProArgSerProGlnArgArgTh 323
   |||||
1004 TGTTCGCGCTG.....GTTGC.....GCCGAG 1026
   |||||
323 rGlyThrValGlnProAlaArgCysProTPrProHisArgAlaAla 340
   |||||
1027 CCGGCAAAATACCTCATCGACGCGGTACAAACCTCGGCAATTCGAAAA 1076
   |||||
340 laGlyProProArgArgArgGlySerGlyAlaProAlaProLeuGlnArgThr 356
   |||||
1077 CAA.....ACTTTCAGTTTCACACAGCCGTCAGCGCGGACG 1117
   |||||
357 ArgSerPheGlyThrAlaGlyLysAlaHisProTPrProAlaArgArgPr 373
   |||||
1118 GCGGCAT..... 1124
   |||||
373 oglyHisTPr***SerAlaAlaAlaAlaProAlaThrGlyAlaProAlaC 390
   |||||
1125 .....GGTGCATGTGTACTTACGAGCGCGGATCGCTTGATATCCT 1169
   |||||
390 yArgAlaGlySerTPrValSerAlaAlaProProAlaGlnGlyArgPro 406
   |||||
1170 GCC.....CACCT.....GCTTTGCGCGATT 1192
   |||||
407 AlaArgAlaArgArgHisProGlyArgCysProGlnAlaSerGlyProAr 423
   |||||
1193 TATTCGTGCGCA.....TACGACAGCGCGGAGGC..... 1223
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146 GlyLeuGlyGly.....AlaGlyPhePr 153
 384 GACTGGCTGCCACCGCTCCGTTCAGCAAAATTCCTGGCGCATGGCG 433
 153 oThrGlyVal.....LysLeuGlnGlyGlyAspL 164
 434 ACCCGTGGCCATCTTCGTAATGCGATGACCAATCCGCTGGCTGCC 483
 164 yAlaGlnThrLeuLeuLeuAlaAlaGluGlyGlyProTyrLeuThr 180
 484 GACCCACGCTCATATCAAGAACGCCGCCGAGATTCAACGCGGCT 533
 181 AlaAspAspArgLeuMetGlnAspCysAlaAlaGlnValGluGlyI 197
 534 GTTGGTATGAGCCGTTGACCGAA..... 558
 197 eArgLeuLeuAlaHisIleLeuGlnProArgGluIleuIleGlyLeu 214
 559CGCAAAATCCATGTTGTAAAGCAGCTGGCGCAGAC 594
 214 LuAspAsnLysProGlnAlaIleSerMetLeuArgAlaValLeuAlaAsp 230
 595 GTGCGCTGAAAATGCTGCCAACATCGAACACATGATTCGGCGGCC 644
 231SerAsnAspIleSerLeuArgValIleProThrLy 242
 645 GCATCTCCCGCT.....TTGAGTGGCAGC 670
 242 sTyrProSerGlyGlyAlaLysGlnLeuThrTyrIleLeuThrGlyLysG 259
 671 ACATTCAATTATGAGCCGCTGGCGGCAATAAACCGCTGGACCAT 720
 259 LysVal.....ProHisGlyArgSerSerAspIleGlyVal 271
 721 AATTATCAAGATGTAATTACC.....ATTGCGGCTGTTGTCAC 761
 272 LeuMetGlnAsnValGlyThrAlaThrAlaValLysArgAlaValIleAs 288
 762 AGCGCGCTGTACACGACGCGCGTATGCCCTAGTGTTGTCACATCA 811
 288 pGlyGluProIleThrGluArgValIleThrLeuThrGlyLysAlaIleAs 305
 812 ACAACGCGCGCTCTGTACGCTGTTGGTGGAAGATGCAAT 861
 305 LaArgProGlyAsnValIleThrAlaArgLeuGlyThrProValArgHisLeu 321
 862ACTGGGCGCAATTGGTTGACACAGACACCGCGTATTCGG 905
 322 LeuAsnAspAlaGlyPheCysProSerAlaAspGlnMetValIleMetG 338
 906 TTCGATTGACGCG 921
 338 yGlyProLeuMetGly 343

seq_name: /SIDSI/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAV04955

seq_documentation_block:

ID AAV04955 standard: Protein: 573 AA.

AC AAV04955;

DT 06-JUL-1999 (first entry)

DE Mycobacterium species protein sequence 41T#3.

KM Secreted protein: Mycobacterium; primer: PCR; amplification: probe;

KW hybridisation: detection; vaccine; immunisation; infection.

OS Mycobacterium sp.

XX W09090186-A2.

PN 25-FEB-1999.

PD

XX
 PF 14-AUG-1998; 98MO-FR01813.
 XX
 PR 11-SEP-1997; 97ER-0011325.
 PR 14-AUG-1997; 97ER-0010404.
 XX
 PA (INSP) INST PASTEUR.
 XX
 PI Gicquel B, Lim EM, Pellicle V, Portnoi D, Goguet de la Salmoniere Y;
 PI Guineno A;
 DR WPI: 1999-181045/15.
 DR N-PSDB: AAX34206.
 XX
 PT Mycobacterial DNA vectors containing reporter constructs - for
 PT Identifying coding or promoter sequences involved in
 PT infection-associated protein expression
 XX
 PS Claim 32; Fig 41T; 309pp; French.
 XX
 CC Sequences AAV04742-Y05000 and AAV07201-Y07204 represent secreted
 CC proteins from various Mycobacterium species microorganisms. The
 CC encoding nucleotide sequences can be used as primers and probes for
 CC methods for detecting and identifying mycobacteria, especially belonging
 CC to the M. tuberculosis complex. The encoded proteins can be used in
 CC vaccines for immunisation against a bacterial or viral infection.
 XX
 SQ Sequence 573 AA;

alignment_scores:

Quality: 114.00 Length: 472
 Ratio: 0.576 Gaps: 24
 Percent Similarity: 41.949 Percent Identity: 25.424

alignment_block:

US-09-303-518D-125 x AAV04955 ..

Align seg 1/1 to: AAV04955 from: 1 to: 573

36 CCGCGGACACCGGAGCAAGCCGTTACGACGCGCCGCCATTCACGAG 85
 162 ArgGlyGlyAlaGlyAsnTyrArgLeuGlyAlaAlaGlyArgArgSer 178
 86 TCGC.....GTTGCTGGCGAAGATATGCGCGGATCGCCCTCGATG 129
 178 rArgArgProValArgAlaArgGlyValGlyArgCysGlyHisArgArg 195
 130 AAGTCAMAGAGGCGGATGCGTCAMAAAGGCCAAGTGTGTTGAGA 179
 195 rg**ArgGlyGlyHisArgAlaGlyLysAspProArgThrAla**Arg 211
 180 CAAMAGAAATCCGGGCGGTGTACTGCGCCGCTTCAGGCAAAATCG 229
 212 AlaArgArgCysGlyArgGly.....GlyArgArg 221
 230 CCGCATTCACCGTGGCGAAGGCGCGTACTTCAGTACGATGATTCGCC 279
 221 gArgThrGlyProAlaGlySerAlaGlyArgValAlaLeuHisHisLeu 238
 280 GTTGAAGCAACGACGAAATGCAATGACGTCACGTCACGTCGAGCGT 329
 238 rGlyAlaGlyThrCysProGlyGlyLeuArgGlyThrLeu.....Ala 250
 330 GGCAAACTTAAGCGCGAGAGAGTGGCGGACCTGATCCAAATCCGGT 379
 251 AlaArgValAlaAspArgHisGlyTyrProThrProArgProAlaIle 267
 380 TGTGGAC.....TGGCTGCGCAGCCGCTCGC 405
 267 gGlyAspValSerValGlyGlyMet**CysCysSerGlyGlyProVal 284
 406 TTCAGCAAAATTCCTGCCGTCGATGCGGACCGTTCGCAATCTTCGCA 455

500 ATATGACCGTA.....GGTCGCGAGCC.....AG 475
 261 ValmetThrIleCysAlaThrProCysTrpSerAlaIaIaGlyAsnLeu 277
 474 CGGATGGGTCATCGCATTCGACGAGATG.....GCGACGCGT 434
 277 rIlySerLeuSerThrCysValThrValMetArgThrIlySerArgGlyS 294
 433 CGGATCGACGCGAGAAATTTGCTGAACGAGCGGTCGCGACGCGATC 384
 294 eRMetAlaThrGlySerLeuAlaValSerGly.....SerSerAlaSer 308
 383 CACAAACCGATTCGATCAGTTGGCGCGCATCTTGCGCG 342
 309 ProSerGlnCysGlyIleArgTrpAlaProThrAlaSerPro 322
 seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1998.DAT:AAW80699
 seq_documentation_block:
 ID AAW80699 standard; Protein: 835 AA.
 AC AAW80699;
 DT 24-DEC-1998 (first entry)
 DE S. pneumoniae cation transporting ATPase.
 XX Streptococcus pneumoniae protein; recombinant; gene expression; DNA chip;
 KM virulence; antibody; infection; detection; treatment; hypothetical;
 KM cell wall biosynthetic; external target; minimal gene set protein.
 XX Streptococcus pneumoniae.
 OS
 PN W09826072-A1.
 XX 18-JUN-1998.
 PD 09-DEC-1997; 97WO-US22578.
 PF 13-DEC-1996; 96US-0036281.
 PR
 XX (ELIL) LILLY & CO ELI.
 PA
 PI Baltz RH, Burgett SG, Dehoff BS, Hoskins JA, Jaskunas SR;
 PI Mills BJ, Norris FH, Peery RB, Rockey PK, Rostreck PR;
 PI Skatrud PL, Smith MC, Solenberg RJ, Treadway RJ;
 PI Young Bellido ML;
 XX WPI: 1998-348529/30.
 DR N-PSDB: AAW65261.
 XX
 PT Streptococcus pneumoniae nucleic acid sequences - used in DNA chips
 PT for evaluating gene expression, and identification of virulence
 PT genes
 PS Claim 3; Pages 288-291; 333pp; English.
 XX
 CC This sequence represents a Streptococcus pneumoniae cation transporting
 CC ATPase. The invention provides DNA sequences (AAW65261 to AAW65304) from
 CC the Streptococcus pneumoniae genome and corresponding protein sequences
 CC (AAW80605 to AAW80728). The protein sequences are classified as
 CC hypothetical, cell wall biosynthetic, external target, or minimal gene
 CC set proteins. A recombinant host containing a vector comprising any of
 CC the above nucleic acids can be used for the recombinant expression of the
 CC proteins. The invention also provides a DNA chip having arrayed on it at
 CC least 15 base pair fragment of any one or more of these DNA sequences.
 CC The DNA chip can be used methods for evaluating gene expression in S.
 CC pneumoniae and for identifying virulence genes in S. pneumoniae.
 CC Antibodies that selectively bind to the above proteins or peptide
 CC fragments can be used to treat S. pneumoniae infection. The antibodies
 CC can also be used to detect S. pneumoniae cells.
 XX
 XX Sequence 835 AA;

alignment_scores:
 Quality: 108.00 Length: 356
 Ratio: 0.617 Gaps: 15
 Percent Similarity: 49.157 Percent Identity: 23.315
 alignment_block:
 US-09-303-518D-125 x AAW80699 ..
 Align seq 1/1 to: AAW80699 from: 1 to: 835
 28 CTGCCCATCCGCGGACAGCCGAGCAAGCCGTTTACGACGCCGCCGCAT 77
 224 LeuAlaValAlaAlaIleProGluGlyLeu.....ProAlaI 236
 78 TACCGAAGTCGCGTTG...CTTGCGCAAGAAATTCGCGGTATCGCCCT 124
 236 eValThrIleValLeuSerLeuGlyThrGlnValLeuAlaLysArgHis 253
 125 CGATG.....AAGTCAGGAAGCGGATGCGTCACAAAGGCGCAAGT 168
 253 eIleValArgGlyLeuProAlaValGluThrLeuGlySerThrGluIle 269
 169 CTGTTTGAGACAAAGAAAT.....CC 191
 270 IleAlaSerAspLysThrGlyThrLeuThrMetAsnLysMetThrValG 286
 192 GGGCGTGCTGTTTCTGCGCGCGCTTCAGCAAAATCGCCGATTCACC 241
 286 uLysValPheThrAspAlaValLeuHisAspSerAlaAspAlaIleuL 303
 242 GTGGC...GAAAGCGGCTACTCAGTCAGTCGATTCGATTCGATGAGC 288
 303 euGlyLeuGluMetProLeuLeuArgSerValValLeuAlaAsnAspThr 319
 289 AAGCAGCAATCGATTGTAAGCTACGACCTGAGCGCTGGCAACTT 338
 320 LysIleAspValGlu.....GlyAsnLe 327
 339 AAGCGCGCAAGAGTGGCGCGCAACCTGATCCAAATCGGTTTGGACTG 388
 327 uIleGlyAspProThrGlnThrAlaPheIleGlnThrAlaLeuAspLysG 344
 389 CGCTCGCACCCGTCGCTTC...AGCAAAATTCCTGCGGTGATGCCGAG 435
 344 LYThrAspValLysGlyPheLeuGluLysTyProArgValAlaGluLeu 360
 436 CCGTTCGCCATCTTGTCAATGCCATGACAC...AATCGGTGGCTGC 482
 361 ProPheAspSerAspArgLysLeuMetSerThrValHisProLeuProAs 377
 483 GAGCCCTACGGTCATTATCAAGAGCCGAGGATTCAAACGCGGCC 532
 377 pSerArgPheLeuValAlaValLysGlyAlaProAsp..... 389
 533 TGTGTGATTGAGCCGTTTGACCGAAGCCAAATCATGTTTGAAGCA 582
 390GlnLeuLeuLysArgCysLeuLeuArgAspLysAla 401
 583 GCTGGCGCGAGAGTCGCGCTGAAATGCTGCCAAC...ATCGAAAGCA 629
 402 GlyAspIleAlaProIleAspGluValThrAsnLeuIleHisThrAs 418
 630 TGAATTGCGCGCGCATCTGC...GGTTGATGGGACGACCA 673
 418 nAsnSerGluMetAlaHisGlnAlaLeuArgValLeuAlaGlyAlaTy 434
 674 TTCATTATTCAGCGCGGTGCGCGGAATAAACCGTGGACCATCAT 723
 435 ..LysIleIleAspSerIleProGluAsnLeuThrSerGluLeuGlu 450
 724 TATCAAGATGTAATTACCATTCGCGGTTTGTTCACAGCGCGTCTGAA 773

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451 ...AspAspLeuIlePheThrGlyLeuIle.....GlyMetIleAs 463
774 CACCGAGCGCGTATGATCCCTAGTGGTTCCTACGACCAACACCGCGCC 823
463 pProGluArgProGluAlaGluAlaValAlaArgValAlaLysGluAlaG 480
824 TCCTGGCGTACCGTTTGGTGGCGAAGATGCGCAATTAATCTGCGCGCA 873
480 LylLeuArgProIleMetIleThrGlyAspHisGlnAspThrAlaGluAla 496
874 TTGGTT.....GACACGAGAACACCGCGT 896
497 IleAlaLysArgLeuGlyIleIleAspAlaAsnAspThrGlnGlyHisVa 513
897 GATTTCGGTGGTGGTATTAAGACGCGCGATGACACAGCGCGCGACGATT 946
513 IleuThrGlyAlaGluLeuAsnGlnLeuSerAspGluGlnGlyLysV 530
947 ATTTGGGACGCTACCAACAATCAGATTCCGTTATGCAAGACCGCGACG 996
530 aValGlyGlnIleThrSerValIleAlaArgValSerProGluHisLysVal 546
997 AAAGAGCTGTGGCGTGG 1014
547 ArgIleValLysAlaIleTrp 552

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seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:ABG05146

seq_documentation_block:

ID ABG05146 standard; Protein; 440 AA.

AC ABG05146;

DT 13-FEB-2002 (first entry)

DE Novel human diagnostic protein #5137.

KW Human: chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder.

OS Homo sapiens.

PN W0200175067-A2.

PD 11-OCT-2001.

PF 30-MAR-2001; 2001WO-US08631.

PR 31-MAR-2000; 2000US-0540217.

PR 23-AUG-2000; 2000US-0649167.

PA (HYSE-) HYSEQ INC.

PI Drmanac RT, Liu C, Tang YT;

DR WPI; 2001-639362/73.

DR N-PSDB; AAS69333.

PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity

PS Claim 20; SEQ ID NO 35505; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome
XX and gene mapping, and in recombinant production of (II). The
XX polynucleotides are also used in diagnostics as expressed sequence tags
XX for identifying expressed genes. (I) is useful in gene therapy techniques
XX to restore normal activity of (II) or to treat disease states involving

CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. ABG00010-ABG30377 represent novel human
CC diagnostic amino acid sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 440 AA;

alignment_scores:

Quality:	106.50	Length:	489
Ratio:	0.598	Gaps:	24
Percent Similarity:	36.401	Percent Identity:	20.654

alignment_block:

US-09-303-518D-125 x ABG05146 ..

Align seg 1/1 to: ABG05146 from: 1 to: 440

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32 CCATTCGGGGGAGACCGGAGCAACCGTTACGACG..... 67
   ||| ||||| ||||| |||||
50 ProArgArgAlaAlaThrProGlnProGlnThrProProSerSerVa 66
68 .....GCCCGGCCATACGA 83
66 LglnProCysAlaAlaThrProCysAlaArgLysGlyAlaProAlaProA 83
84 AGTCGGCTGCTGGCGAAGAAATATGCGGTATGCGCCCTGATGAAG 133
   ||| ||||| ||||| |||||
83 IacIValAlaThrTrpProThrThrArgProSerMetAla**ArgThrSer 99
134 TCAGGAGAGCGG.....ATGCCGTCAAAAAGGCCAAGTGTCTG 171
   ||| ||||| ||||| |||||
100 ***AlaSerThrThrAlaProArgMetAlaProProArgSerHisAlaCy 116
172 TTGAAGACAAAAGAAATCCGGGCGGTGTACTGCGCGCGCTTCAGG 221
   ||| ||||| ||||| |||||
116 sAlaArgSerHisArgProGluThr.....AlaArgSerAlaA 129
222 CAAATTCGGCGCGATTCACCGTGGCGAAAGCGCTACTTCAGTCAGTCG 271
   ||| ||||| ||||| |||||
129 rGThrAlaProArgSerAlaIleThrArgArgAlaPheThrSerThrArg 145
272 TGATTGGCGCTGAAGCAACGCAAAATCGATTGAACGCTACGACCT 321
   ||| ||||| ||||| |||||
146 ProProProThrThrArgThrValAlaSerSer.....GlyThrHisTh 160
322 GAAGCGCTGGCAAACTTAAGCGCGGAGAGAGTGGCGCAACTGATGCA 371
   ||| ||||| ||||| |||||
160 rSerGlyLeuSerProThrAlaSerArgLeuAla..... 171
372 ATCCGGTTTGTGATCGGCTGC...GCACCGTCCGTTACGCAAAATTC 418
   ||| ||||| ||||| |||||
172 .....ArgCysArgAlaProGlyArgSerSerThrIle 182
419 CTG..... 421
183 IleThr***ThrCysArgSerProThrArgLeuCysCysProAlaGlnAr 199
422 ..CGTCGATGCCGAGCCGTCGCAATCTTCGTCAATGCA..... 460
199 gProLeuProAlaSerSerProSerSerSerArgThrSerArgSerV 216
461 ..TGCACACCAATCCGCTGGCTGGCGACCTTACGCTATTTCAAAGAG 508

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Align seg 1/1 to: ABG23389 from: 1 to: 1194

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1133 TTGCGCAACAGCGGGCGGCTATTCGATTGTCGCCGGGAGACAGACCTGCA 1264
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
584 LeuSerGlnSerGlyProProGlyLeuLeuPro..... 594
1263 CAAGAGGAGGCTCTTCGTCGCAATTCAGACCAACCAATGCC...TCG 1217
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
595 .....SerProSerPheAspSerIysProProThrThrLeuLeuG 608
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
1216 CGCTGCGGTATCGCGGAGATTAATGCGCAAGACAGGCTGGCGCAGG 1167
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
608 LylleuIleProAlaProSerMet..... 615
1166 ATATCAAGGGGATCAGCGCGCTCGTAAGTACCAATCGCACCATGGCGCG 1117
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
616 .....ValPro..AlaThrAspThrIly 622
1116 TTGCGCGCGGCTTGAGCGCTGTTGACTGAGAGATTGTTTTCAGGA 1067
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
622 SalAProProThrLeuGlnAlaGlnThrAlaThrLysProGlnAlaThrS 639
1066 AATGCCGAGGAGTTGTACGCGGTGATGAGATTTGTCGGGTGGCGGCA 1017
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
639 erAlaProSerProAlaProLysGlnSerPheLeuPheGlyThrGlnAsn 655
1016 ACCAAGCGCAAGAGCTCTTTCGCGCATTCCTTCGATTAACGAAATCTG 967
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
656 ThrSerProSerSer.....ProAlaAla..... 663
966 ATTGTGTAGAGCGCCCAATATATGTCGCGCTTGTGTATGCGCGCT 917
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
664 .....ProAlaAlaSerSerAlaProMetPheLysProI b76
916 TCAATACCGAAGCGGAAATCAGCGGTTGTGTGTCAACCAATGCGCC 867
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
676 LephThrAlaPro..... 680
866 GCATATTTGCGTACTTTCGCAACCAACGATGCGCAGAGCGCGG 817
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
681 .....ProLysSer.....GlnLysGlnI 687
816 TTGTGTACTTGAGAACCACTAGGCAATCAGCGCGGTGTGACAG 767
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
687 yProThr.....ProProGlyProSerValThrAlaThrAla..... 699
766 GGGCTGTGCAACCAAGCGCAATGATTAATCATCTTGATTAATGATG 717
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
700 ..ProSerSerSerLeuProThrThr..... 708
716 GTCCACAGCGTTTATTCGCGCGGCGGCTGATGAATGAATGCGCT 667
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
709 .....ThrSerThrThrAlaProThr..... 715
666 GCCACTCAAAACG.....GCAGGATGCGGCGCGGCAATTCATGTGTT 623
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
716 ....PheGlnProValPheSerSerMetGlyProProAlaSer...ValP 730
622 CGATGTGGCAGCATTTTCAGACGCGAGCTGTGCGCAGCTCCCTTACAA 573
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
730 roleuProAlaProPhePheLysGlnThrThrProAlaThrAlaPro 746
572 ACATGATTTTGTGCGTCAACGCGCTCATACCAACAGCGCGCTT 523
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
747 Thr..... 747
522 GAAATCTGCGCGCTCTTTGATATGACCGTAGGCTGGCAGCGCAG 473
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
748 .....ThrThrAlaProLeuPheThrGlyLeuAlaSerAlaThrSera 762
472 GATTGGTGCCATGCAATGACAGAGATGGCAACGCGCTCGCATCG... 426
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

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762 laValAlaProIleThrSerAlaSerProSerThrAspSerAlaSerLys 778
425 ...ACGGCAGGAATTTTCTGAACGAGCGGTGCGAGCGACCATCCACAA 379
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
779 ProAlaPheGlyPheGlyIleAsnSer...ValSerSerSerValSe 794
378 ACCGGATTTGATCAGGTTGGCGGCACCTTCTTCCCGCTTAAGTTGCCA 329
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
794 rThrThrThrSerThrAlaThrAlaThrAlaSerGlnProPheLeuPheGlyA 811
328 GC.....GCTTCAGGTGCGTACGCTCAAACTCGAATTCGTCGTG... 288
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
811 laProGlnAlaSerAlaAlaSerPheThrProAlaMetGlySerIlePhe 827
287 .....CCTTCACGCGCAATCAGCACTGACTGAAGTACGCGCTT 250
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
828 GlnPheGlyLysProProAlaLeuProThrThrThrValThrThrPh 844
249 TTGCGCAGGTGAATCGCGGATTTTGGCTGAAGCGCGCGCATTAACA 200
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
844 eSerGlnSer...LeuHisThrAlaValAlaProThrAlaThrSerSera 860
199 CCAGC.....CCCGGATCTTTTGTCTTCAACAGCACTGGCGCTTT 156
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
860 laAlaAspPheSerGlyPheGlySerThrLeuAlaThrSerAlaProAla 876
155 TTGACGGCATCGCCTTCTTCTGACTTCATCGAGGCGCATACGGCAT 106
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
877 ThrSerSerGlnProThrLeuThrPheSerAsnThrSerThrProThrPh 893
105 TTCTTCGCCCAACGACGACCTTCGTAATGAGCGCGGCGCGTAAACG 56
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
893 eAsnIleProPheGlySerSerAlaLysSerProLeuProSerIyrProg 910
55 CTTCGCTCCGCTGCGCGCGATGGC 30
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
910 lAlaAsnProGlnProAlaPheGly 918
seq_name: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1197.DAT:AAW13504
seq_documentation_block:
ID AAW13504 standard; protein; 1644 AA.
XX
AC AAW13504;
XX
DT 30-JAN-1998 (first entry)
XX
DE B. bronchiseptica adenylcyclase-haemolysin mutant delta-Cla.
XX
KW Mutant; Bordetella pertussis; adenylcyclase; haemolysin; wild type;
KW deletion; induction; antibody; Bordetella parapertussis; vaccine; human;
KW Bordetella bronchiseptica; veterinary.
XX
OS Bordetella bronchiseptica.
XX
FH Synthetic.
XX
PH Key
XX
FT Region 826..827
FT Location/Qualifiers
FT /note= "deletion of amino acids 827-887 from the wild
FT type sequence between these positions"
FT Modified-site 919..924
FT /note= "Arg at position 922 is modified by addition of
FT a fatty acid"
XX
FT FT
XX
XX FR2736064-A1.
XX
XX PD 03-JAN-1997.
XX
XX PD 30-JUN-1995; 95FR-0007945.
XX
XX PR 30-JUN-1995; 95FR-0007945.
XX
XX PA (INSP ) INST PASTEUR.

```



```
XX AC AAY04954;
XX DT 06-JUL-1999 (first entry)
XX DE Mycobacterium species protein sequence 41IT#2.
XX KW Secreted protein; Mycobacterium; primer; PCR; amplification; probe;
XX KM hybridisation; detection; vaccine; immunisation; infection.
XX OS Mycobacterium sp.
XX PN WC0909186-A2.
XX PD 25-FEB-1999.
XX PE 14-AUG-1998; 98WO-FR01813.
XX PR 11-SEP-1997; 97FR-0011325.
XX PR 14-AUG-1997; 97FR-0010404.
XX PA (INSP ) INST PASTEUR.
XX PI Glacquel B, Lim EM, Pelicic V, Portnoi D, Goguet de la Salmoniere Y,
XX PI Guigueno A;
XX DR WPI: 1999-181045/15.
XX DR N-PSDB: AAX34206.
XX PT Mycobacterial DNA vectors containing reporter constructs - for
XX PT identifying coding or promoter sequences involved in
XX PT infection-associated protein expression
XX PS Claim 32; Fig 41T; 309pp; French.
XX CC Sequences AAY04742-Y05000 and AAY07201-Y07204 represent secreted
XX CC proteins from various Mycobacterium species microorganisms. The
XX CC encoding nucleotide sequences can be used as primers and probes for
XX CC methods for detecting and identifying mycobacteria, especially belong
XX CC to the M. tuberculosis complex. The encoded proteins can be used in
XX CC vaccines for immunisation against a bacterial or viral infection.
XX SQ Sequence 572 AA;

alignment_scores:
    Quality: 106.00      Length: 509
    Ratio: 0.586        Gaps: 33
    Percent Similarity: 35.560      Percent Identity: 20.825

alignment_block:
US-09-303-518D-125 x AAY04954 ..

Align seg 1/1 to: AAY04954 from: 1 to: 572

26 ACCTGCCATCGCGGACGACCGGACCGTTACGACGCGCGCC 75
   :::::::::::::::::::: |||
113 SerCysProArgSerCysAlaIysSerGlnArg***ProArgLeuArgPr 129
   :::::::::::::::::::: |||
76 ATTACCGAGTCCGCTTGGTGGCGAAGATATGCGGTA...TGGCGC 122
   ||| :::::::::::::::::::: |||
129 oProProLeuAlaArgTyrCysGlyArgSerThrProThrProSerGlyP 146
   ||| :::::::::::::::::::: |||
123 CTGATGAAGAATCGAAGAGCGCATCGCTCAAAAAG..... 160
   ||| :::::::::::::::::::: |||
146 roArgCysGlyAlaAlaSerSerThrProThrProValAlaArg 162
   ||| :::::::::::::::::::: |||
161 .....GCCAAGTGTCT.....TT 174
   :::::::::::::::::::: |||
163 ArgCysArgGluLeuSerSerArgCysGlyProProGluProse 179
   :::::::::::::::::::: |||
175 GAAGAAGAAAGATCCGGCGTGGTTACTGCGCGGCTTCAGGCA 224
   :::::::::::::::::::: |||
```

```
179 rThrsAlaArgThrArgGlyTrp.....ProVal 190
225 AATGCCCGCATTCACCGTGGCGAAGACCGTACTTCACTCAGTCGTA 274
   :::::::::::::::::::: |||
190 rSerProProProValThrArgArgSerSerArgTrpLysArg..... 204
   ||| :::::::::::::::::::: |||
275 TTGGCGTTGAAGGACGACGAAATCGATTTCAGCTTCAGCAGCTGAA 324
   ||| :::::::::::::::::::: |||
205 ...ProProAsnSerLeuThrCysSerProMetArgAla..... 216
   ||| :::::::::::::::::::: |||
325 GCGCTGGCAACCTTAAAGCGGGAAGAGTGGCGCGCAACCTGATCCATC 374
   ||| :::::::::::::::::::: |||
217 ArgTrp...ThrProAlaSerGlyAlaCysTrp..... 226
   ||| :::::::::::::::::::: |||
375 CGGTTGTGAGCTGCGTGGCGACCGCTCCGTTTCAGCAAAATTCCTCGC 424
   ||| :::::::::::::::::::: |||
227 ...PheCysTrpThrArgCysAlaProProSerAlaGlyArgHisLeuPro 242
   ||| :::::::::::::::::::: |||
425 TCGATGCGGAGCCGCTTGCATCTTCATGATGATGAGCAATCCG 474
   ||| :::::::::::::::::::: |||
243 GlyArg...SerThrAsnProArgArgAla...ArgCysArgProThrAr 257
   ||| :::::::::::::::::::: |||
475 CTGCTGCGGACCCCTACGGTCAATTATCAAGAGCCGCGAGATTTCAA 524
   ||| :::::::::::::::::::: |||
257 9.....LeuProAsnAlaProProArgAsnSerA 267
   ||| :::::::::::::::::::: |||
525 ACGCGGCG...TGTGTATGTAGCGCGTTGACCGCAACCAAAATCATG 571
   ||| :::::::::::::::::::: |||
267 rG***CysIleCysTrp..... 272
   ||| :::::::::::::::::::: |||
572 TTTGTAGGCGAGTGGCGGACGACGTCGCGCTGAATAATGTCGAACATC 621
   ||| :::::::::::::::::::: |||
273 .....ArgTyrValMetLeu.GlnArgA 280
   ||| :::::::::::::::::::: |||
622 GAACACATGAATTGGCGGCGCGCCGATCTCGCGGTTTGAG..... 662
   ||| :::::::::::::::::::: |||
280 rG.....ThrsCysGlyIleAspSerArgAsn 289
   ||| :::::::::::::::::::: |||
663 TGGACGACATTCATTTCATGACGCGGTGGCGCAATAAACCGGT 712
   |||
290 Trp..... 290
713 GGACCATCATATTATCAGATGTAATTACCATGCGCGTTGTTCACA 762
   :::::::::::::::::::: |||
291 .....ValSerArg.....TrpProSerProLeuAla 300
   ||| :::::::::::::::::::: |||
763 GCGCGTGAACACGACGCGCGGTGATGCGCT..... 794
   ||| :::::::::::::::::::: |||
300 rGProThrAlaThrProTyrThrSerThrProThrThrProValProPro 316
   ||| :::::::::::::::::::: |||
795 .....AGGTGGTTCATCAATCAACAAACCGCGCTCTTGC 829
   ||| :::::::::::::::::::: |||
317 TrpLysProAspTrpArgTrp.GlyGluLeuAlaGlySer***SerArgA 333
   ||| :::::::::::::::::::: |||
830 GTACCGTTTGGGTGCGAAGATATCGCAAAATTACTGCGGCGAATTGGTT 879
   ||| :::::::::::::::::::: |||
333 rGSerValProGly..... 337
   ||| :::::::::::::::::::: |||
880 GACACAGACAAACCGCGTATTCGCGTTCGATTTGAAGCGCGGATTC 929
   :::::::::::::::::::: |||
338 ProAlaAspCysArgProValAlaGly..... 346
   ||| :::::::::::::::::::: |||
930 ACAAGCGCGACGATATTGAGACCTACCAATCAGATTTCGCTTA 979
   :::::::::::::::::::: |||
347 ArgGlyAlaAlaProCysTrpArgSerSerThrAlaThrValPro... 361
   ||| :::::::::::::::::::: |||
980 TCGAAGAGAGCGCGACGAAGAGCTGTTCGCTGGGTGGCGCGCAC... 1027
   ||| :::::::::::::::::::: |||
362 .....ProSerCysSerProGlyArgAlaProAlaLacy 372
   ||| :::::::::::::::::::: |||
1028 .....CGGCAATATCATCAGCGGTACCAACCTCGCGCATTTCT 1070
   ||| :::::::::::::::::::: |||
372 scYAspArgValAlaInThrPro**HisArgProProIleSerVal.... 387
```



```

632 AATTCGGCGCGCCGATCTCCGCTTGTAGTGGACGACATTCATTTTC 681
171 ....GlyArgAlaAlaArgAsnArgHisLeuSerAlaHisSer...A 185
682 ATGACCGCGGTGGCGGAAATAAACCGTGGACCATTAATTCACAGA 731
185 rgrArgAlaLeuGlnGlyProGlnGlyArgAspAlaGlyIle... 199
732 TGTAAATACCATTGGCGCTTGTGTTCAC... 761
200 ....LeuTyrLeuProAspLeuSerArgAspArgAlaLeuArgGln 214
762 ....AGCGG...TCGAACACCGACCGCGGTATGCCCTAGGTGCT 804
214 gLeuArgProGlnGlyArgHisProAlaValAlaValLeuGlnGlyLeu 231
805 CAAGTCACAAACCGCGCTTGGCTACCGCTTGGGTC... 845
231 rArgArgGlnGlnIleProAlaAspGlnProHisGlyGlyGlyThr 247
846 .....GAAAGTATCGCAATTAATCTCGGCGCAATTG 877
248 ArgArgAspGlyThrPheGlnAspPheArgAlaTrpArgGlnValAl 264
878 TTGACACAGACAAACCGCGGTATGCCGTGCTATGAA... 917
264 a.....ArgLeuProArgAsnLeuPro...GlyValGlnArgSerVal 278
918 .....CGGCGGATTCACAAAGCGCGACGATTTATGGGACGCTA 959
278 IsrProHisArgArgArgHisGlnArgGlnGlyArg...GlyArgLeu 292
960 CCACATACGATTCCTGTTATCGAAGAGCGCGACAAAGCGCTGTTCG 1009
293 GlnArgHisAspArgArgHisProArgGlySerGln... 304
1010 GCTGGGTTGGCGCGCGACCAATCTCCACAC... 1046
305 .....AlaAspProHisGlyArgAlaLeuHisArgLeuProLeuPhe 319
1047 .....GCGTACAAACCTCGCGCATTCCTGTA 1073
319 InGlnGlnProArgGlnAspAlaAlaLeuGlyPheArgValHisProArg 335
1074 AAACAAATCTTCAAGTTCACACACGCGTCACGCGCGCGCGCA 1123
336 ArgThrArgLeuSerValAsp.....GlnAlaArgArgAspArgHis 349
1124 TGGTGGCGATGTACTTACGAGCGCGTATGCCCTTGATATCTGCC 1173
349 sGlyAlaAspGlnGlyIleArg.....GlnValG 359
1174 ACCCTGCTTTGGCGGATTAACTGTCGCGGATACGACGCGCGAGC 1223
359 InPro.....GlnArgGlnArgArgHisArgArgGly 369
1224 ATTGGGTGCTTGAATTGCA.....CGAAGAAAGACCTGCTT 1261
370 SerProGlnLeuGlnArgAlaValAlaHisValAlaArgHisProArg 386
1262 TGTGCA.....CTTCGCTGCTCCG 1281
386 rProGlnAspAlaAlaLysAsnProPheGlyArgTrpArgLeuAsnLeuPro 403
1282 GGCAAATACGAATAGCGCGCTGTGCGCAAGTCTGGAAC 1325
403 Ly.....GlnAlaAlaGlyAsn 408
seq_name: /std1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT.AAB59824
seq_documentation_block:
ID AAB59824 standard; Protein: 1605 AA.
XX

```

```

AC AAB59824:
XX
XX 04-APR-2001 (first entry)
XX
XX Protein #1 encoded by TufD/E gene.
XX
XX DE
XX
XX KM Toluene degradation; enzyme: waste degradation; TufD.
XX
XX OS Thauera aromatica.
XX OS Xanthomonas maltophilia.
XX OS Geobacter metallireducens.
XX OS Azoriscus toluyticus.
XX
XX PN WO200072650-A2.
XX
XX PD 07-DEC-2000.
XX
XX PF 24-MAY-2000; 2000WO-US14298.
XX
XX PR 01-JUN-1999; 99US-0323872.
XX
XX PA (UOH-) UNITV OHIO.
XX
XX PI Coschigano PW;
XX
XX DR WPI: 2001-041080/05.
XX
XX N-PSDB: AAF23627.
XX
XX PT Composition comprising toluene degrading enzyme useful for biological
XX treatment of organic compounds, especially for degrading toluene or its
XX analogs
XX
XX PS Disclosure: Fig 12; 122pp; English.
XX
XX CC The present invention relates to toluene degrading enzyme genes and
XX proteins tufH (see AAF23629 and AAB59831), tufI (AAF23630 and AAB59832),
XX tufR (AAF23631 and AAB59833) and tufG (AAF23632 and AAB59834). The
XX toluene degrading enzymes are homologues of pyruvate formate lyase. The
XX CC toluene degrading enzymes are useful for biological treatment of organic
XX compounds and in particular for the degradation of toluene and its
XX CC analogs contained in liquid or solid waste source. The present sequence
XX is a protein sequence encoded by toluene degrading enzyme gene, TufD/E.
XX
XX SQ Sequence 1605 AA;

```

```

alignment_scores:
Quality: 106.00 Length: 498
Ratio: 0.535 Gaps: 27
Percent Similarity: 39.759 Percent Identity: 24.096

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alignment_block:

US-09-303-518d-125 x AAB59824 ..

Align seg 1/1 to: AAB59824 from: 1 to: 1605

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18 AGCTTAAACCTGCGCCATCGGGCGACAGCCGAGCAACCCGTTA..... 62
603 ArgThrValProHisGlyArgAlaGlyLeuProAlaGlnArgLeuAl 619
63 .....CGACGCGCGCGCATTAACCGAAGTCGCTTGGCGAA 102
619 AlaAlaGlyArgArgGlyGly.....GlyAsp 629
103 GAATATGCGCGGTATGCGCCCTCGATGAAGTCAAGAGGCGATGCCGT 152
629 InLeuGlnAlaAlaProAla...GlnGlnValSerAlaLeuPheArg 644
153 CAAAAAGGCCAAGTGTGTAAGACAAATAATCCGCGCGCTGTGT 202
645 SerGlyArgProArgProHisValSerGlyGlnGlnHisGlyAlaVal 661
203 TTAAGTGGCGCGCTTCAAGGCAAAATCC.....CGCATTCACCGT 243

```


CC discloses genomic DNA sequences (AB116176-AB130511), expressed DNA sequences (AB101840-AB116175) and the encoded proteins

CC (AB57737-AB572072).

CC The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 1908 AA:

Alignment_scores:

Quality: 105.50 Length: 557
Ratio: 0.498 Gaps: 31
Percent Similarity: 38.061 Percent Identity: 21.544

Alignment_block:

US-09-303-518D-125 x ABB70137 ..

Align seg 1/1 to: ABB70137 from: 1 to: 1908

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29 TCGCCCA...TCGCGGCGACAGCCGAGCAAGCCGTTTACGAGCGCCCGCC 75
||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
859 CysProvalGlnArgGlnArgSerProSerCysLysArgThrAsnPr 875
76 ATTACCGAAGTCGCGCTGCTGGCGAAGATATGCGCGGTATCGCCCTC 125
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
875 GlnAsnArgSerArg.....LysGlnProProIleCysSerAsn 889
126 GATGAAGATCAAGAGGCGATCGCGTCAAAAAAGCCCAAGTCG..... 169
:: :||| :||| :||| :||| :||| :||| :||| :||| :|||
889 IuThrLeuMetGlnLysProValLysSerTyArgPheSerCysProGlu 905
170 .....TGTTCAGACAAAGAAAGATCCGCGCGTGTGTTACTGCG 210
||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
906 AsnGlyLysCysAlaSerCysArgLysLeuLysProLysCysHisLysAl 922
211 CCGGTTAGCGCAAAATCGCCGCGATTCACCGTGGCGAAAGCCGCTACT 260
||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
922 aLeuAlaGlnLysLys..... 927
261 TCAGTCAGTCGATGATTCGCTTGAAGCGACAGCAAAATCGATTTGAC 310
927 ..... 927
311 GCTACGCACTGAAGCGCTGCGCAAACTTAACGCGGAGAAAGTGGCGCG 360
:|||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
928 .....LysThrSerSerAlaLysLysCysAlaAl 937
361 AACCTGATCCAAAT.....CGGTTTGTGACTGCGCTGCGCGACCCG 401
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
937 aSerValAlaGlnLysCysCysProMetCysGlyLeu..... 949
402 TCCGTTAGCAAAATTCCTGCGCTGATGCGGAGCCGCTTCG...CGATC 448
||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
950 .....LeuProMetMetProAsnIleSerIleProAsp 960
449 TCGTCATG.....CGATGACACC 468
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
961 ValGlySerGlyAsnArgIleIleTyLysThrLeuGlyArgCysArgPr 977
469 AATCCGCTGGCTGCGCGCCTACGG..... 493
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
977 O...ArgLysLeuProLysThrArgTyGlyLeuThrIleCysCysLysL 993
493 ..... 493
993 ysArgCysGlnGlyArgAsnSerHisTyMetThrGlyThrValAla 1009
494 ...TCATATCAAGAAG..... 508
||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1010 ThrSerLeuLysArgAlaLysAlaAlaMetThrGlyValAsnProVa 1026
509 .....CCGCCGAGATTTCAAAACGCGCGCTGTGATTTGAGCGCTTGAC 554
```

```
||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1026 AlaProProArgValArgGlnSer.....P 1035
555 CG.....ACGCAAAATCCATGTTTGTAAAGCAGCTGGCGCAG 592
|| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1035 rOpheProProLeuAsnMetProSerHisProValSerGln..... 1048
593 ACGTGGCGCTGAAATGCTGCCAACATGCACATGATTCGGCGCG 642
||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1049 .....ProThrGly...HisValArgSerAla 1057
643 CCGCATCTCGCGGTTTGAAGTGGACAGACATTCATTTATGAGCGCGT 692
|||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1057 rArg.....AlaGlyHisSerTyLeuAspGlnAspArgS 1069
693 CGGCG.....CGATTAACCCGTGTGACCATCA 721
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1069 eSerProSerThrSerAlaGlnProArgGlnArgProAlaProPro 1085
722 ATTATCAAGATTAATTACCATTCGCGCTTGTTCACACAG..... 764
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1086 ProPheLysMetArgSerProSerAlaAsnValAlaGln.LysSerGluL 1102
765 .....CCGTCGAACACCGGAGCGGTGAT 788
||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1102 ysArgGlnGlyGlyTyArgLnisProSerGlnMetArgThrArgThr 1118
789 TGCCCT.....AGTGTTCCTCAAGTCAACAACCCGCTTCCTGC 829
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1119 MetGlnProValIleGlnIleHisSerThrProLysGlnAlaIleThrAl 1135
830 GTACCGCTTTGGGTGGCAAGATATCGCAATTAATCTCGGCGCAATTGGTT 879
||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1135 aTyArg..... 1137
880 GACACAGACACCGCGTATTCGCGTGTGATTAAGCGCGCATTAAC 929
||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1138 .....SerGlnProGluProGluProSerAsnArgSerGlnArgGluPro 1152
930 ACNAGCGCGCACAGATTATTGGAGCGCTACACAAATCAATATTCGCTTA 979
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1153 ThrAsnArgSerGln.....ThrGluProProAlaSer.....Se 1164
980 TCGAAGACGCGCGCAG..... 995
||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1164 rArgSerArgProGluThrProAspLeuProThrSerLeuIleAlaProA 1181
996 .....CAAGAGCTGTTCGCGCTGCGCGCGCA 1025
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1181 lAserMetAspAlaGlnIleThrGlnIleProValArgArgGlyAlaSerTy 1197
1026 GCGCGCAAAATCTCCATCAGCGCTACCAACCTCGCGCATTTCTCGAAA 1075
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1198 LeuGlyThrGlnLeuAsn.....ProPheTyArgG1 1208
1076 ACAACCTTCAAGTTCAACAAGCGCTCAACGCGCGCGCGCCAGT 1125
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1208 ySerPheLeuArgValArgHisSerArgAspSerProGlnSerArg.Leu 1224
1126 GTGCGGATGTGACTTACGACGCGGTGATGCGCTGATATCTGCCAC 1175
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1225 SerProIleHisGlnArgGlyGlnIleSerGlnValAlaProArgProThr 1241
1176 CTTGCTTTGCGCGCATTAATCGTGGCGGATACC.....GACAGCGCGC 1219
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1241 rValProGluMetGlyLeuProValLysTyThrArgLeuSerGlnLeuG 1258
1220 AGGATTTGGTT..... 1231
||||| :||| :||| :||| :||| :||| :||| :||| :||| :|||
1258 lAlaA.ThrValrPheGlyAspProPheValLysGlyLysThrAsnThrG1 1274
1232 .GCTTGAATGG.....ACGACAGACCTCGCTTGTG 1265
| :||| :||| :||| :||| :||| :||| :||| :||| :|||
```

1274 yThrtPserTrpArgGlnIlePheGlyArgLysLysSerProThrT 1291
 1266 CA.....GCTTCGTCTGCCCGGCAATACGATACGGCCGC 1303
 :::::::::::|||||:::
 1291 hrLysAspLysProAlaLysThrMetLaspAlaAsnArgArgMetGlyGly 1307
 1304 TGTTCGCGCAAGTGC 1318
 |||
 1308 AsnCysProIleCys 1312
 |||
 seq_name: /SIDSI/gcgdata/geneseq/geneseqp-emb1/AA2001.DAT:ABG00972
 seq_documentation_block:
 ID ABG00972 standard; Protein; 4274 AA.
 XX
 AC ABG00972;
 XX
 DT 13-FEB-2002 (first entry)
 XX
 DE Novel human diagnostic protein #963.
 XX
 KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
 food supplement; medical imaging; diagnostic; genetic disorder.
 XX
 OS Homo sapiens.
 XX
 FN WO200175067-A2.
 XX
 PD 11-OCT-2001.
 XX
 PF 30-MAR-2001; 2001WO-US08631.
 XX
 PR 31-MAR-2000; 2000US-0540217.
 PR 23-AUG-2000; 2000US-0649167.
 XX
 PA (HYSE-) HYSEQ INC.
 XX
 PI Drmanac RT, Liu C, Tang YT;
 XX
 DR WPI: 2001-639362/73.
 DR N-PSDB: AAS65159.
 XX
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 PS
 PS Claim 20; SEQ ID NO 31331; 103bp; English.
 XX
 CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for creating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. ABG00010-ABG30377 represent novel human
 CC diagnostic amino acid sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 4274 AA;

alignment_scores:
 Quality: 105.50 Length: 454
 Ratio: 0.533 Gaps: 17
 Percent Similarity: 43.612 Percent Identity: 21.366
 alignment_block:
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 1212 1212
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 692ACCGGCTGATGAA 679
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649 GATGC.....GGGCGCGCGAAT 633
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1751 SerValSerSerValValSerAlaAlaThrAspThrValGluLysValPh 1767
596 .....ACGT 593
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1811 ThrSerSerValThrSerSerIleIleThrValProValLysSerVal 1828
458 .....GCATTGACGAAGATG.....GCCACGCGC 435
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1845 PheThrLysSerAlaAlaAlaLeuLeuSerProIleLysThrIleuThr 1861
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1861 rGluThrHisProGlnProHisPheSerArgThrSerSerProValLys 1878
334 TTGCC.....AGCGTTTCAGGTCGTCAGGTCGAACACTGATTCGTCG 291
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290 TTGCCTTCACAG 279
1895 LeuSerSerSer 1898

seq_name: /sids1/gcgdata/geneseq/geneseqp-emb1/AA2001.DAT:ABG07375
seq_documentation_block:
ID ABG07375 Standard; Protein; 4386 AA.
XX
AC ABG07375;
XX
XX 13-FEB-2002 (first entry)
XX
DE Novel human diagnostic protein #7366.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.

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PR 23-AUG-2000; 2000US-0649167.
XX
XX (HYSE-) HYSEQ INC.
XX
XX Drmanac RT, Liu C, Tang YT;
XX
XX WPI: 2001-639362/73.
XX
XX N-PSDB; AAS71562.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity
PS
PS Claim 20; SEQ ID No 37734; 103bp; English.
XX
XX The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome
XX and gene mapping, and in recombinant production of (II). The
XX polynucleotides are also used in diagnostics as expressed sequence tags
XX for identifying expressed genes. (I) is useful in gene therapy techniques
XX to restore normal activity of (II) or to treat disease states involving
XX (II). (II) is useful for generating antibodies against it, detecting or
XX quantitating a polypeptide in tissue, as molecular weight markers and as
XX a food supplement. (II) and its binding partners are useful in medical
XX imaging of sites expressing (II). (I) and (II) are useful for treating
XX disorders involving aberrant protein expression or biological activity.
XX The polypeptide and polynucleotide sequences have applications in
XX diagnostics, forensics, gene mapping, identification of mutations
XX and to produce other types of data and products dependent on DNA and
XX amino acid sequences. ABG00010-ABG30377 represent novel human
XX diagnostic amino acid sequences of the invention.
XX Note: The sequence data for this patent did not appear in the printed
XX specification, but was obtained in electronic format directly from WIPO
XX at ftp.wipo.int/pub/published_pcl_sequences.
XX
XX Sequence 4386 AA:

alignment_scores:
Quality: 105.50 Length: 454
Ratio: 0.533 Gaps: 17
Percent Similarity: 43.612 Percent Identity: 21.366

alignment_block:
US-09-303-518d-125/rev x ABG07375 ..
Align seg 1/1 to: ABG07375 from: 1 to: 4386

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1243 CCAATTCACAGCAACCCATGCTTCGCGCGTG..... 1212
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1212 ..... 1212
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